



# **STIC Search Report**

## **Biotech-Chem Library**

**STIC Database Tracking Number: 168245**

**TO: Jeffrey Fredman**  
**Location: rem/2C89/2C18**  
**Art Unit: 1637**  
**Friday, October 14, 2005**

**Case Serial Number: 09/744097**

**From: Edward Hart**  
**Location: Biotech-Chem Library**  
**REM-1A55**  
**Phone: 571-272-2512**

**edward.hart@uspto.gov**

### **Search Notes**

Examiner Fredman,

Here are the results of the search you requested.

Please feel free to contact me if you have any questions.

Edward Hart

STIC-Biotech/ChemLib

168245

mg

From: Fredman, Jeffrey  
Sent: Tuesday, October 11, 2005 10:23 AM  
To: STIC-Biotech/ChemLib  
Subject: 09/744,097

RECEIVED  
OCT 11 2005  
STIC-BIOTECH/CHM LIB  
(STIC)

Please search SEQ ID NO: 76 in nucleic acid databases.

Thanks,

Jeffrey Fredman  
Art Unit 1637  
Remsen Building 2C89  
(571)272-0742

2C18

\*\*\*\*\*

Searcher: \_\_\_\_\_  
Searcher Phone: \_\_\_\_\_  
Date Searcher Picked up: 10/12/05  
Date completed: 10/14/05  
Searcher Prep Time: \_\_\_\_\_  
Online Time: \_\_\_\_\_

\*\*\*\*\*

Type of Search  
NA# 1 AA# \_\_\_\_\_  
S/L: \_\_\_\_\_ Oligomer: \_\_\_\_\_  
Encode/Transl: \_\_\_\_\_  
Structure #: \_\_\_\_\_ Text: \_\_\_\_\_  
Inventor: \_\_\_\_\_ Litigation: \_\_\_\_\_

\*\*\*\*\*

Vendors and cost where applicable  
STN: \_\_\_\_\_  
DIALOG: \_\_\_\_\_  
QUESTEL/ORBIT: \_\_\_\_\_  
LEXIS/NEXIS: \_\_\_\_\_  
SEQUENCE SYSTEM: \_\_\_\_\_  
WWW/Internet: \_\_\_\_\_  
Other (Specify): \_\_\_\_\_

GenCore version 5.1.6  
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: October 13, 2005, 17:46:38 ; Search time 1479 Seconds  
(without alignments)  
884.578 Million cell updates/sec

Title: US-09-744-097A-76  
Sequence: 1 gtacgctagctaccctaggtctaggc 27

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 4708233 seqs, 24227607955 residues

Total number of hits satisfying chosen parameters: 9416466

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

GenEmbl.\*

- 1: gb.ba.\*
- 2: gb.htg.\*
- 3: gb.in.\*
- 4: gb.om.\*
- 5: gb.ov.\*
- 6: gb.pat.\*
- 7: gb.ph.\*
- 8: gb.pl.\*
- 9: gb.pr.\*
- 10: gb.ro.\*
- 11: gb.sts.\*
- 12: gb.sy.\*
- 13: gb.un.\*
- 14: gb.vl.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	20.6	76.3	135412	2	AC148279
2	20.6	76.3	156661	2	AC148358
3	20.6	76.3	278310	2	AC127955
C 4	20.6	76.3	285193	2	AC111242
C 5	20.6	76.3	293962	2	AC112303
6	20.4	75.6	188946	10	AC127236
C 7	20.4	75.6	194613	2	AC116089
C 8	20.2	74.8	124104	8	AC134931
9	19.8	73.3	119743	2	AC134515
10	19.8	73.3	211465	2	AC103070
11	19.8	73.3	212559	2	AC095896
12	19.6	72.6	857	4	SSAPOA11
13	19.6	72.6	5551	10	AK173314
C 14	19.6	72.6	207223	10	AL732521
15	19.6	72.6	212481	10	AC126959
16	19.6	72.6	252851	2	AC135094
C 17	19.2	71.1	276137	2	AC131170
18	19.2	71.1	91977	10	AL732588
19	19.2	71.1	116926	10	AL808012

20	19.2	71.1	147467	2	AC118573
C 21	19.2	71.1	186449	2	AC147134
22	19.2	71.1	188892	2	AC091327
C 23	19.2	71.1	237378	2	AC087038
24	19	70.4	110000	1	AE017180.27
25	19	70.4	150222	2	AC120016
26	19	70.4	164530	10	AL596104
27	19	70.4	187514	2	AC109240
C 28	19	70.4	203946	2	AC069465
C 29	19	70.4	233852	2	AC108549
C 30	19	70.4	235241	10	AL928893
C 31	19	70.4	240264	2	AC107434
32	19	70.4	245134	2	AC126639
33	19	70.4	248917	2	AC111455
C 34	19	70.4	250579	2	AC126843
35	19	70.4	325223	2	AC109504
C 36	18.8	69.6	81368	2	AC094244.4
C 37	18.8	69.6	103428	2	AC096436.6
C 38	18.8	69.6	104113	10	AL928640
C 39	18.8	69.6	110000	2	AC102028.3
40	18.8	69.6	142937	2	AC139916
C 41	18.8	69.6	148320	10	AL669849
C 42	18.8	69.6	154685	10	AL606511
43	18.8	69.6	157088	10	AC127292
C 44	18.8	69.6	171936	10	AL929035
C 45	18.8	69.6	173598	10	AL672194

## ALIGNMENTS

AC148279 135412 bp DNA linear HTG 15-APR-2004  
Sorex araneus clone SA\_Ba-546N4, WORKING DRAFT SEQUENCE, 7 ordered pieces.

AC148279

AC148279.2 GI:46391177

HTG; HTGS PHASE2; HTGS\_DRAFT.

Sorex araneus (European shrew)

Sorex araneus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Insectivora; Soricidae; Soricinae; Sorex.

1 (bases 1 to 135412)

Antonsellis,A., Avelle,K., Benjamin,B., Blakesley,R.W.

Bouffard,G.G., Brinkley,C., Brooks,S., Chu,G., Coleman,B.,

Coleman,H., Daki,N., Engle,J., Granite,S., Guan,X., Gupta,J.,

Haghighi,P., Han,J., Hansen,N., Ho,S.-L., Hu,P., Hurle,B.,

Idol,J.R., Jones,C., Karlins,E., Kim,H., Kwong,P., Laric,P.,

Larson,S., Lee-Lin,S.-O., Legaspi,R., Maduro,O.L., Maduro,V.B.,

Margulies,E.H., Masiello,C., Maskeri,B., McDowell,J.,

Mullikin,J.C., Paguirigan,C., Portnoy,M.E., Prased,A., Puri,O.,

Reddix-Dugue,N., Schandler,K., Schueler,M.G., Shah,K., Sison,C.,

Santripop,S., Thomas,J.W., Thomas,P.J., Tsipouri,V., Vogt,J.L.,

Wetherby,K.D., Young,A. and Green,E.D.

NISC Comparative Sequencing Initiative

Unpublished

2 (bases 1 to 135412)

Green,E.D.

Direct Submission

Submitted (19-FEB-2004) NIH Intramural Sequencing Center, 8717

Grovenomt Circle, Gaithersburg, MD 20877, USA

3 (bases 1 to 135412)

Green,E.D.

Direct Submission

Submitted (15-APR-2004) NIH Intramural Sequencing Center, 8717

Grovenomt Circle, Gaithersburg, MD 20877, USA

On Apr 15, 2004 this sequence version replaced gi:42627935.

----- Genome Center

Center: NIH Intramural Sequencing Center

Center code: NISC

Web site: <http://www.nisc.nih.gov>

Contact: [nisc\\_zoo@nhgri.nih.gov](mailto:nisc_zoo@nhgri.nih.gov)



coverage in Q20 bases and has been reviewed to rule out gross misassemblies, the low-quality ends of sequence contigs have been trimmed away, and each base is associated with a Phrap-derived quality score.

----- Summary Statistics

Sequencing vector: plasmid; n/a; 100% of reads  
 Chemistry: Dye-terminator Big Dye; 100% of reads  
 Assembly program: Phrap; version 0.990319  
 Consensus quality: 154585 bases at least Q40  
 Consensus quality: 155145 bases at least Q30  
 Consensus quality: 155506 bases at least Q20  
 Insert size: 107000; agarose-fp  
 Insert size: 155661; sum-of-contigs  
 Quality coverage: 12.02x in Q20 bases; agarose-fp  
 Quality coverage: 8.26x in Q20 bases; sum-of-contigs

-----  
 \* NOTE: This is a 'working draft' sequence. It currently consists of 11 contigs. Gaps between the contigs are represented as runs of N. The order of the pieces is believed to be correct as given, however the sizes of the gaps between them are based on estimates that have been provided by the submitter.

\* This sequence will be replaced  
 \* by the finished sequence as soon as it is available and  
 \* the accession number will be preserved.

\* 1 41408: contig of 41408 bp in length  
 \* 41409: gap of 2126 bp in length  
 \* 43634: contig of 2126 bp in length  
 \* 43635: gap of unknown length  
 \* 43735: contig of 13527 bp in length  
 \* 57262: gap of unknown length  
 \* 57361: contig of 6500 bp in length  
 \* 63861: gap of unknown length  
 \* 63862: contig of 33808 bp in length  
 \* 63962: gap of unknown length  
 \* 97770: contig of 10730 bp in length  
 \* 97870: contig of 10730 bp in length  
 \* 108699: gap of unknown length  
 \* 109700: contig of 20478 bp in length  
 \* 129178: contig of 5413 bp in length  
 \* 129278: gap of unknown length  
 \* 134691: contig of 7141 bp in length  
 \* 134791: gap of unknown length  
 \* 141931: contig of 9880 bp in length  
 \* 142032: gap of unknown length  
 \* 151911: contig of 9880 bp in length  
 \* 151912: gap of unknown length  
 \* 152012: contig of 4650 bp in length.

FEATURES

source

1. 155661  
 /organism="Sorex araneus"  
 /mol\_type="genomic DNA"  
 /db\_xref="taxon:42254"  
 /clone="SA\_Ba-521C10"  
 /clone\_lib="SA\_Ba"  
 1. 67579

/note="clone overlaps with GenBank Accession Number AC148279, clone SA\_Ba-546N4 (center project name gbi)"

1. 41408  
 /note="assembly\_fragment"  
 clone\_end:SP6  
 vector\_side:left

41509. 43634  
 /note="assembly\_fragment"

43735. 57261  
 /note="assembly\_fragment"

57362. 63861  
 /note="assembly\_fragment"

63962. 97769  
 /note="assembly\_fragment"

97870. 108599  
 /note="assembly\_fragment"

108700. 129177  
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129278. 134690  
 /note="assembly\_fragment"

misc\_feature  
 129878. 156661  
 /note="clone overlaps with GenBank Accession Number AC148355, clone SA\_Ba-408O18 (center project name gga)"  
 134791. 141931  
 /note="assembly\_fragment"  
 142032. 151911  
 /note="assembly\_fragment"  
 152012. 156661  
 /note="assembly\_fragment"  
 clone\_end:T7  
 vector\_side:right

ORIGIN

Query Match 76.3%; Score 20.6; DB 2; Length 156661;  
 Best Local Similarity 85.2%; Fred. No. 20;  
 Matches 23; Conservative 0; Mismatches 4; Indels 0; Gaps 0;  
 Oy 1 GTAGCTAGCTACCCCTAGGTCTAGGC 27  
 Db 39469 GTACCTACTACCCCTAGATTAGGC 39495

RESULT 3

AC127955

LOCUS

DEFINITION Rattus norvegicus clone CH230-270L18, \*\*\* SEQUENCING IN PROGRESS  
 AC127955 278310 bp DNA linear HTG 19-SEP-2002  
 \*\*\*, 6 unordered pieces.

ACCESSION

AC127955.2 GI:23195969

VERSION HTG; HTGS\_PHASE1; HTGS\_DRAFT; HTGS\_ENRICHED.

KEYWORDS Rattus norvegicus (Norway rat)

SOURCE Rattus norvegicus

ORGANISM Rattus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;

Rattus.

1 (bases 1 to 278310)

REFERENCE

AUTHORS

Muzny, D. Marie., Metzker, M. Lee., Abramson, S., Adams, C., Alder, J., Allen, C., Allen, H., Alebrooks, S., Amin, A., Anguiano, D., Anyalebechi, V., Aoyagi, A., Ayodeji, M., Baca, E., Baden, H., Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Benahmed, F., Biswal, K., Blair, J., Blankenburg, K., Blyth, P., Brown, M., Bryant, N., Buhay, C., Burch, P., Burrell, K., Calderon, E., Cardenas, V., Carter, K., Cavazos, I., Ceasar, H., Center, A., Chacko, J., Chavez, D., Chen, G., Chen, R., Chen, Y., Chen, Z., Chu, J., Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Cree, A., D'Souza, L., Davila, M. L., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D., Delgado, O., Denson, S., Deramo, C., Ding, Y., Dinh, H., Divya, K., Draper, H., Dugan-Rocha, S., Dunn, A., Durbin, K., Duval, B., Eaves, K., Egan, A., Escotto, M., Eugene, C., Evans, C. A., Falls, T., Fan, G., Fernandez, S., Finley, M., Flagg, N., Forbes, L., Foster, M., Foster, P., Fraser, C. M., Gabisi, A., Ganta, R., Garcia, A., Garner, I., Garza, M., Gebregeorgis, E., Geer, K., Gill, R., Grady, M., Guerra, W., Guevara, W., Gunaratne, P., Haaland, W., Hamill, C., Hamilton, C., Hamilton, K., Harvey, Y., Havlak, P., Hawes, A., Henderson, N., Hernandez, J., Hernandez, R., Hines, S., Hladun, S. L., Hodgson, A., Hogues, M., Hollins, B., Howells, S., Hulyk, S., Hume, J., Idlebird, D., Jackson, A., Jackson, L., Jacob, L., Jiang, H., Johnson, B., Johnson, R., Jolivet, A., Karpathy, S., Kelly, S., Khan, Z., King, L., Kovar, C., Kowis, C., Kraft, C. L., Lebow, H., Levan, J., Lewis, L., Li, Z., Liu, J., Liu, J., Liu, W., Liu, Y., London, P., Longacre, S., Lopez, J., Lorensuhewa, L., Loulseghe, H., Lozano, R. J., Lu, X., Ma, J., Maheshwari, M., Mahindartne, M., Mahmoud, M., Malloy, K., Mangum, A., Mangum, B., Mapua, P., Martin, K., Martin, R., Martinez, E., Mawhinney, S., McLeod, M. P., McNeill, T. Z., Meenen, E., Milosavljevic, A., Miner, G., Minja, E., Montemayor, J., Moore, S., Morgan, M., Morris, K., Morris, S., Munidasa, M., Murphy, M., Nair, L., Nankervis, C., Neal, D., Newton, N., Nguyen, N., Norris, S., Nwaokemele, O., Okwuonu, G., Olarnpunsagoon, A., Pal, S., Parks, K., Pasternak, S., Paul, H., Perez, A., Perez, B., Pfankoch, C., Plopper, F., Poindexter, A., Popovic, D., Primus, E., Pu, L. L., Puazo, M., Quiroz, J., Rachlin, E., Reeves, K., Regier, M. A., Reigh, R., Reilly, B., Reilly, M., Ren, Y., Reuter, M., Richards, S., Riggs, F.,

Rives, C., Rodkey, T., Rojas, A., Rose, M., Rose, R., Ruiz, S.J.,  
 Sanders, W., Savery, G., Scherer, S., Scott, G., Shatsman, S., Shen, H.,  
 Shetty, J., Shvartsbeyn, A., Sison, I., Sitter, C.D., Smajls, D.,  
 Sneed, A., Sodergren, E., Song, X.-Z., Sorrelle, R., Sosa, J.,  
 Steimle, M., Strong, R., Sutton, A., Svatek, A., Tabor, P., Taylor, C.,  
 Taylor, T., Thomas, N., Thomas, S., Tingey, A., Trejos, Z., Umani, K.,  
 Valas, R., Vera, V., Villalana, D., Waldron, L., Walker, B., Wang, J.,  
 Wang, Q., Wang, S., Warren, J., Warren, R., Wei, X., White, F.,  
 Williams, G., Willson, R., Wleciyk, R., Wooden, H., Worley, K.,  
 Wright, D., Wright, R., Wu, J., Yakub, S., Yen, J., Yoon, V.,  
 Yu, F., Zhang, J., Zhou, X., Zhou, X., Zhao, S., Dunn, D., von  
 Niederhausern, A., Weiss, R., Smith, D.R., Holt, R.A., Smith, H.O.,  
 Weinstock, G. and Gibbs, R.A.

**TITLE**  
**JOURNAL**  
**REFERENCE**  
**AUTHORS**  
**JOURNAL**

Unpublished  
 2 (bases 1 to 278310)  
 Worley K.C.

**TITLE**  
**JOURNAL**

Submitted (19-JUL-2002) Human Genome Sequencing Center, Department  
 of Molecular and Human Genetics, Baylor College of Medicine, One  
 Baylor Plaza, Houston, TX 77030, USA

3 (bases 1 to 278310)  
 Rat Genome Sequencing Consortium.

**REFERENCE**  
**AUTHORS**  
**JOURNAL**

Submitted (19-SEP-2002) Human Genome Sequencing Center, Department  
 of Molecular and Human Genetics, Baylor College of Medicine, One  
 Baylor Plaza, Houston, TX 77030, USA

On Sep 19, 2002 this sequence version replaced gi:21908490.  
 The sequence in this assembly is a combination of BAC based reads  
 and whole genome shotgun sequencing reads assembled using Atlas  
 (http://www.hgsc.bcm.tmc.edu/projects/rat/). As a result, the  
 sequence may extend beyond the ends of the clone and there may be  
 contigs that consist entirely of whole genome shotgun sequence  
 reads. Both end sequences and whole genome shotgun sequence only  
 contigs will be indicated in the feature table.

----- Genome Center -----  
 Center: Baylor College of Medicine  
 Center code: BCM  
 Web site: http://www.hgsc.bcm.tmc.edu/  
 Contact: hgsc-help@bcm.tmc.edu  
 ----- Project Information -----  
 Center project name: KBR  
 Center clone name: CH230-270L18  
 ----- Summary Statistics -----  
 Assembly program: Phrap; version 0.990329  
 Consensus quality: 159405 bases at least Q40  
 Consensus quality: 163261 bases at least Q30  
 Consensus quality: 165346 bases at least Q20  
 Estimated insert size: 206482; sum-of-contigs estimation  
 Quality coverage: 3x in Q20 bases; sum-of-contigs estimation

----- NOTE: Estimated insert size may differ from sequence length  
 (see http://www.hgsc.bcm.tmc.edu/docs/Genbank\_draft\_data.html).  
 \* NOTE: This is a 'working draft' sequence. It currently  
 \* consists of 6 contigs. The true order of the pieces  
 \* is not known and their order in this sequence record is  
 \* arbitrary. Gaps between the contigs are represented as  
 \* runs of N, but the exact sizes of the gaps are unknown.  
 \* This record will be updated with the finished sequence  
 \* as soon as it is available and the accession number will  
 \* be preserved.

1 197344: contig of 197344 bp in length  
 \* 197345: 197444: gap of unknown length  
 \* 197445 266948: contig of 69504 bp in length  
 \* 266949 267048: gap of unknown length  
 \* 267049 267114: contig of 2666 bp in length  
 \* 267115 269814: gap of unknown length  
 \* 269815 271799: contig of 1985 bp in length  
 \* 271800 271899: gap of unknown length  
 \* 271900 273966: contig of 2067 bp in length  
 \* 273967 274066: gap of unknown length  
 \* 274067 278310: contig of 4244 bp in length.

Location/Qualifiers

source  
 1. 278310  
 /organism="Rattus norvegicus"  
 /mol\_type="genomic DNA"  
 /db\_xref="taxon:10116"  
 /clone="CH230-270L18"  
 misc\_feature  
 1. 2376  
 /note="wgs\_contig"  
 misc\_feature  
 153418..156139  
 /note="wgs\_contig"  
 misc\_feature  
 211774..212782  
 /note="wgs\_contig"  
 ORIGIN

Query Match 76.3%; Score 20.6; DB 2; Length 278310;  
 Best Local Similarity 85.2%; Pred. No. 19;  
 Matches 23; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Oy 1 GTACCTAGTACCCCTAGGCTAGGC 27  
 |||||  
 Db 248352 GTAGCAAGCTACCTAGGCTAGC 248378  
 |||||

RESULT 4  
 AC111242/c  
 LOCUS  
 DEFINITION AC111242 285193 bp DNA linear HTG 13-MAY-2003  
 Rattus norvegicus clone CH230-232H4, \*\*\* SEQUENCING IN PROGRESS  
 \*\*\*, 11 unordered pieces.  
 AC111242  
 AC111242.4 GI:30578456  
 HTG: HTGS PHASE3: HTGS DRAFT: HTGS\_ENRICHED.  
 KEYWORDS  
 Rattus norvegicus (Norway rat)  
 SOURCE  
 Rattus norvegicus  
 ORGANISM  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;  
 Rattus.

REFERENCE  
 1 (bases 1 to 285193)  
 Muzny, D., Marie, E., Metzker, M., Lee, S., Adams, C., Alder, J.,  
 Allen, C., Allen, H., Alsbrooks, S., Amin, A., Anguiano, D.,  
 Anyalebechi, V., Ayagi, A., Ayodeji, M., Baca, E., Baden, H.,  
 Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Benahmed, F.,  
 Biswal, K., Blair, J., Blankenburg, K., Blyth, P., Brown, M.,  
 Bryant, N., Buhay, C., Burch, P., Burrell, K., Calderon, E.,  
 Cardenas, V., Carter, K., Cavazos, I., Ceasar, H., Center, A.,  
 Chacko, J., Chavez, D., Chen, G., Chen, R., Chen, Y., Chen, Z., Chu, J.,  
 Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Cree, A., D'Souza, L.,  
 Davila, M.L., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D.,  
 Delgado, O., Denison, S., Deramo, C., Ding, Y., Dinh, H., Divya, K.,  
 Draper, H., Dugan-Rocha, S., Dunn, A., Durbin, K., Duval, B., Eaves, K.,  
 Egan, A., Escotto, M., Eugene, C., Evans, C.A., Falls, T., Fan, G.,  
 Fernandez, S., Finley, M., Flagg, N., Forbes, L., Foster, M., Foster, P.,  
 Fraser, C.M., Gabisi, A., Ganta, R., Garcia, A., Garner, T., Garza, M.,  
 Gebregeorgis, E., Geer, K., Gill, R., Grady, M., Guerra, W., Guevara, W.,  
 Gunaratne, P., Haaland, W., Hamill, C., Hamilton, C., Hamilton, J.,  
 Harvey, Y., Havlak, P., Hawes, A., Henderson, N., Hernandez, J.,  
 Hernandez, R., Hines, S., Hladun, S.L., Hodgson, A., Hogues, M.,  
 Hollins, B., Howells, S., Hulyk, S., Hume, J., Idlebird, D., Jackson, A.,  
 Jackson, L., Jacob, L., Jiang, H., Johnson, B., Johnson, R., Jolivet, A.,  
 Karpathy, S., Kelly, S., Kelly, S., Khan, Z., King, L., Kovar, C.,  
 Kowitz, C., Kraft, C.L., Lebow, H., Levan, J., Lewis, L., Li, Z., Liu, J.,  
 Liu, J., Liu, W., Liu, Y., London, P., Longacre, S., Lopez, J.,  
 Lorenschew, L., Loulseghe, H., Lozano, R.J., Lu, X., Ma, J.,  
 Maheshwari, M., Mahindartne, M., Mahmoud, M., Malloy, K., Mangum, A.,  
 Mangum, B., Mapus, P., Martin, K., Martin, R., Martinez, E.,  
 Mawhinney, S., Mcleod, M.P., McNeill, T.Z., Meenen, E.,  
 Milosavljevic, A., Miner, G., Minja, E., Montemayor, J., Moore, S.,  
 Morgan, M., Morris, K., Morris, S., Munidasa, M., Murphy, M., Nair, L.,  
 Nankervis, C., Neal, D., Newton, N., Nguyen, N., Norris, S.,  
 Nwaokeme, O., Okwuonu, G., Olarnpunsagoon, A., Pal, S., Parks, K.,  
 Pasternak, S., Paul, H., Perez, A., Perez, I., Pfannkuch, C.,  
 Pioppo, F., Poindecker, A., Popovic, D., Primus, E., Pu, L.,  
 Puazo, M., Quiroz, J., Rachlin, E., Reeves, K., Regier, M.A., Reigh, R.,  
 Reilly, B., Reilly, M., Ren, Y., Reuter, M., Richards, S., Riggs, F.,  
 Rives, C., Rodkey, T., Rojas, A., Rose, M., Rose, R., Ruiz, S.J.,

Sanders, W., Saverly, G., Scherer, S., Scott, G., Shatsman, S., Shen, H., Shetty, A., Shvartsbeyn, A., Sisson, I., Sitter, C.D., Smajis, D., Speed, A., Sodergren, E., Song, X.-Z., Sorelle, R., Sosa, J., Steimle, M., Strong, R., Sutton, A., Svatek, A., Taber, P., Taylor, C., Taylor, T., Thomas, N., Thomas, S., Tingey, A., Trejos, Z., Umani, K., Valas, R., Vera, V., Villasana, D., Waldron, L., Walker, B., Wang, J., Wang, Q., Wang, S., Warren, J., Warren, R., Wei, X., White, F., Williams, G., Willson, R., Wleczky, R., Wooden, H., Worley, K., Wright, D., Wright, R., Wu, J., Yakub, S., Yen, J., Yoon, L., Yoon, V., Yu, F., Zhang, J., Zhou, J., Zhou, X., Zhao, S., Dunn, D., von Niederhausern, A., Weiss, R., Smith, D.R., Holt, R.A., Smith, H.O., Weinstock, G. and Gibbs, R.A.

# TITLE JOURNAL

## REFERENCE AUTHORS

## TITLE JOURNAL

## REFERENCE AUTHORS

## TITLE JOURNAL

## REFERENCE AUTHORS

## TITLE JOURNAL

## COMMENT

2 (bases 1 to 285193)  
Worley, K.C.

Submitted (19-FEB-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA

3 (bases 1 to 285193)

Rat Genome Sequencing Consortium.

Submitted (13-MAY-2003) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA

On May 13, 2003 this sequence version replaced gi:23604121.

The sequence in this assembly is a combination of BAC based reads and whole genome shotgun sequencing reads assembled using Atlas (<http://www.hgsc.bcm.tmc.edu/projects/rat/>). Each contig described in the feature table below represents a scaffold in the Atlas assembly (a 'contig-scaffold'). Within each contig-scaffold, individual sequence contigs are ordered and oriented, and separated by sized gaps filled with Ns to the estimated size. The sequence may extend beyond the ends of the clone and there may be sequence contigs within a contig-scaffold that consist entirely of whole genome shotgun sequence reads. Both end sequences and whole genome shotgun sequence only contigs will be indicated in the feature table.

----- Genome Center

Center: Baylor College of Medicine

Center code: BCM

Web site: <http://www.hgsc.bcm.tmc.edu/>

Contact: [hgsc-help@bcm.tmc.edu](mailto:hgsc-help@bcm.tmc.edu)

----- Project Information

Center project name: GWE

Center clone name: CH230-232H4

----- Summary Statistics

Assembly program: Atlas 3.0;

Consensus quality: 242831 bases at least Q40

Consensus quality: 249455 bases at least Q30

Consensus quality: 253771 bases at least Q20

Estimated insert size: 272024; sum-of-contigs estimation

Quality coverage: 5x in Q20 bases; sum-of-contigs estimation

-----

\* NOTE: Estimated insert size may differ from sequence length

(see [http://www.hgsc.bcm.tmc.edu/docs/Genbank\\_draft\\_data.html](http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html))

\* NOTE: This sequence may represent more than one clone.

\* NOTE: This is a 'working draft' sequence. It currently consists of 11 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown.

\* This record will be updated with the finished sequence, as soon as it is available and the accession number will be preserved.

1 5183: contig of 5183 bp in length

5184 5283: gap of unknown length

5284 17822: contig of 12539 bp in length

17823 17822: gap of unknown length

116644: contig of 98722 bp in length

116645 116744: gap of unknown length

116745 157368: contig of 40624 bp in length

\* 157369 157468: gap of unknown length

\* 157469 247503: contig of 90435 bp in length

\* 247504 248003: gap of unknown length

\* 248004 261891: contig of 13888 bp in length

\* 261892 261991: gap of unknown length

\* 261992 276510: contig of 14519 bp in length

\* 276511 276610: gap of unknown length

\* 276611 277878: contig of 1268 bp in length

\* 277879 277978: gap of unknown length

\* 277979 279325: contig of 1347 bp in length

\* 279326 279425: gap of unknown length

\* 279426 282175: contig of 2750 bp in length

\* 282176 282275: gap of unknown length

\* 282276 285193: contig of 2918 bp in length.

# FEATURES

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/organism="Rattus norvegicus"  
/mol\_type="genomic DNA"  
/db\_xref="taxon:10116"  
/clone="CH230-232H4"

## misc\_feature

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clone\_end:T7  
complement(7161..8031)  
/note="clone\_boundary"  
clone\_end:T7  
site:EcoRI

## misc\_feature

end\_sequence:BZ092926"  
16581..17822  
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34961..39614  
/note="wgs\_contig"  
40916..42387  
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58872..59764  
/note="clone\_boundary"  
clone\_end:Sp6  
site:EcoRI

## misc\_feature

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clone\_end:Sp6"  
136932..138724  
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clone\_end:Sp6"  
155552..157368  
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clone\_end:Sp6"  
257732..259507  
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clone\_end:Sp6"  
268633..274031  
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clone\_end:Sp6"

## misc\_feature

Query Match 76.3%; Score 20.6; DB 2; Length 285193;  
Best Local Similarity 85.2%; Pred. No. 19;  
Matches 23; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

## misc\_feature

1 GTAGCCTAGCTACCCCTAGGCTTAGGC 27  
|||||

## misc\_feature

Db 114184 GTAGCCAGCTACCTCTAGGTGTAGAC 114158  
|||||

## misc\_feature

RESULT 5  
AC112303/c AC112303 293962 bp DNA linear HTG 15-NOV-2002  
LOCUS Rattus norvegicus clone CH230-208p20, \*\*\* SEQUENCING IN PROGRESS  
DEFINITION \*\*\* 8 unordered pieces.  
ACCESSION AC112303

## misc\_feature

ORIGIN

Query Match 76.3%; Score 20.6; DB 2; Length 285193;  
Best Local Similarity 85.2%; Pred. No. 19;  
Matches 23; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

## misc\_feature

1 GTAGCCTAGCTACCCCTAGGCTTAGGC 27  
|||||

## misc\_feature

Db 114184 GTAGCCAGCTACCTCTAGGTGTAGAC 114158  
|||||

## misc\_feature

RESULT 5  
AC112303/c AC112303 293962 bp DNA linear HTG 15-NOV-2002  
LOCUS Rattus norvegicus clone CH230-208p20, \*\*\* SEQUENCING IN PROGRESS  
DEFINITION \*\*\* 8 unordered pieces.  
ACCESSION AC112303

AC12303.4 GI:24635626  
 HTG; HTGS\_PHASE1; HTGS\_DRAFT; HTGS\_ENRICHED.  
 Rattus norvegicus (Norway rat)  
 ORGANISM  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;  
 Rattus.  
 1 (bases 1 to 293962)  
 Muzny, D., Metzker, M., Lee, A., Abramson, S., Adams, C., Alder, J.,  
 Allen, C., Allen, H., Alsbrooks, S., Amin, A., Anguiano, D.,  
 Anyalebechi, V., Aoyagi, M., Ayodeji, M., Baca, E., Baden, H.,  
 Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Benahmed, F.,  
 Biswal, K., Blair, J., Blankenburg, K., Blyth, P., Brown, M.,  
 Bryant, N., Buhay, J., Burch, P., Burrell, K., Calderon, E.,  
 Cardenas, V., Carter, K., Cavazos, I., Ceasar, H., Center, A.,  
 Chacko, J., Chavez, D., Chen, G., Chen, R., Chen, Y., Chen, J., Chu, J.,  
 Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Cree, A., D'Souza, L.,  
 Davila, M., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D.,  
 Delgado, O., Denson, S., Deramo, C., Ding, X., Dinh, H., Divya, K.,  
 Draper, H., Dugan-Rocha, S., Dunn, A., Durbin, K., Duval, B., Eaves, K.,  
 Egan, A., Escotto, M., Eugene, C., Evans, C.A., Falls, T., Fan, G.,  
 Fernandez, S., Finley, M., Flagg, N., Forbes, L., Foster, M., Foster, P.,  
 Fraser, C.M., Gabisi, A., Ganta, R., Garcia, A., Garner, T., Garza, M.,  
 Gebregorgis, E., Geer, K., Gill, R., Grady, M., Guerra, W., Guevara, W.,  
 Gunaratne, P., Haaland, W., Hamil, C., Hamilton, C., Hamilton, K.,  
 Harvey, Y., Havlak, P., Hawes, A., Henderson, N., Hernandez, J.,  
 Hernandez, R., Hines, S., Hladun, S.L., Hodgson, A., Hognes, M.,  
 Hollins, B., Howells, S., Huiyk, S., Hume, J., Idlebird, D., Jackson, A.,  
 Jackson, L., Jacob, L., Jiang, H., Johnson, B., Johnson, R., Jolivet, A.,  
 Karpathy, S., Kelly, S., Kelly, S., Khan, Z., King, L., Kovar, C.,  
 Kowis, C., Kraft, C.L., Lebow, H., Levan, J., Lewis, L., Li, Z., Liu, J.,  
 Liu, J., Liu, W., Liu, J., London, P., Longacre, S., Lopez, J.,  
 Lorensheva, L., Loulseghe, H., Lozano, R.J., Lu, X., Ma, J.,  
 Maheshwari, M., Mahindartne, M., Mahmoud, M., Malloy, K., Mangum, A.,  
 Mangum, B., Mapa, P., Martin, K., Martin, R., Martinez, E.,  
 Mauney, S., McLeod, M.P., McNeill, T.Z., Meenen, E.,  
 Milosavljevic, A., Miner, G., Minja, E., Montemayor, J., Moore, S.,  
 Morgan, M., Morris, K., Morris, S., Munidasa, M., Murphy, M., Nair, L.,  
 Nankervis, C., Neal, D., Newton, N., Nguyen, N., Norris, S.,  
 Nwankwelu, O., Okwuonu, G., Olarpunsaogon, A., Pal, S., Parks, K.,  
 Pasternak, S., Paul, H., Perez, A., Perez, L., Pfankuch, C.,  
 Plopper, F., Poindexter, A., Popovic, D., Primus, E., Pu, L.-L.,  
 Puazo, M., Quiroz, J., Rachlin, E., Reeves, K., Regier, M.A., Reigh, R.,  
 Reilly, B., Reilly, M., Ren, Y., Reuter, M., Richards, S., Riggs, F.,  
 Rives, C., Rodkey, T., Rojas, A., Rose, M., Rose, R., Ruiz, S.O.,  
 Sanders, W., Savary, G., Scherer, S., Scott, G., Shatman, S., Shen, H.,  
 Shetty, J., Shvartsbeyn, A., Sisson, I., Sitter, C.D., Smajls, D.,  
 Sneed, A., Sodergren, E., Song, X.-Z., Sorelle, R., Sosa, J.,  
 Steimle, M., Strong, R., Sutton, A., Svatek, A., Tabor, P., Taylor, C.,  
 Taylor, T., Thomas, N., Thomas, S., Tingey, A., Trejos, Z., Umani, K.,  
 Valas, R., Vera, V., Villasana, D., Waldron, L., Walker, B., Wang, J.,  
 Wang, Q., Wang, S., Warren, J., Warren, R., Wei, X., White, F.,  
 Williams, G., Willson, R., Wlecyk, R., Woodden, H., Worley, K.,  
 Wright, D., Wright, R., Wu, J., Yakub, S., Yen, J., Yoon, L., Yoon, V.,  
 Yu, F., Zhang, J., Zhou, J., Zhou, X., Zhao, S., Dunn, D., von  
 Niederhausern, A., Weiss, R., Smith, D.R., Holt, R.A., Smith, H.O.,  
 Weinstein, G. and Gibbs, R.A.  
 Direct Submission  
 Unpublished  
 2 (bases 1 to 293962)  
 Worley, K.C.  
 Direct Submission  
 Submitted (21-FEB-2002) Human Genome Sequencing Center, Department  
 of Molecular and Human Genetics, Baylor College of Medicine, One  
 Baylor Plaza, Houston, TX 77030, USA  
 3 (bases 1 to 293962)  
 Rat Genome Sequencing Consortium.  
 Direct Submission  
 Submitted (15-NOV-2002) Human Genome Sequencing Center, Department  
 of Molecular and Human Genetics, Baylor College of Medicine, One  
 Baylor Plaza, Houston, TX 77030, USA  
 On Nov 6, 2002 this sequence version replaced gi:23603916.  
 The sequence in this assembly is a combination of BAC based reads

and whole genome shotgun sequencing reads assembled using Atlas  
 (http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described  
 in the feature table below represents a scaffold in the Atlas  
 assembly (a 'contig-scaffold'). Within each contig-scaffold,  
 individual sequence contigs are ordered and oriented, and separated  
 by sized gaps filled with Ns to the estimated size. The sequence  
 may extend beyond the ends of the clone and there may be sequence  
 contigs within a contig-scaffold that consist entirely of whole  
 genome shotgun sequence reads. Both end sequences and whole genome  
 shotgun sequence only contigs will be indicated in the feature  
 table.

----- Genome Center  
 Center: Baylor College of Medicine  
 Center code: BCM  
 Web site: http://www.hgsc.bcm.tmc.edu/  
 Contact: hgsc-help@bcm.tmc.edu  
 ----- Project Information  
 Center project name: GMYH  
 Center clone name: CH230-208P20  
 ----- Summary Statistics  
 Assembly program: Phrap; version 0.990329  
 Consensus quality: 206616 bases at least Q40  
 Consensus quality: 209134 bases at least Q30  
 Consensus quality: 211305 bases at least Q20  
 Estimated insert size: 210335; sum-of-contigs estimation  
 Quality coverage: 6x in Q20 bases; sum-of-contigs estimation

-----  
 \* NOTE: Estimated insert size may differ from sequence length  
 (see http://www.hgsc.bcm.tmc.edu/docs/Genbank\_draft\_data.html).  
 \* NOTE: This is a 'working draft' sequence. It currently  
 \* consists of 8 contigs. The true order of the pieces  
 \* is not known and their order in this sequence record is  
 \* arbitrary. Gaps between the contigs are represented as  
 \* runs of N, but the exact sizes of the gaps are unknown.  
 \* This record will be updated with the finished sequence  
 \* as soon as it is available and the accession number will  
 \* be preserved.

FEATURES	source	misc_feature	misc_feature	misc_feature	misc_feature	misc_feature
1	27340: contig of 27340 bp in length					
2	27341					
3	27440: gap of unknown length					
4	27441					
5	27440: contig of 24161 bp in length					
6	51601					
7	51701: gap of unknown length					
8	51702					
9	268441: contig of 216740 bp in length					
10	268541: gap of unknown length					
11	268542					
12	270645: contig of 2104 bp in length					
13	270646					
14	270745: gap of unknown length					
15	272746					
16	272238: contig of 1493 bp in length					
17	272239					
18	275335					
19	275354: contig of 3016 bp in length					
20	275455					
21	281008: contig of 5554 bp in length					
22	281009					
23	281108: gap of unknown length					
24	293962: contig of 12854 bp in length.					
25	293962					
26	Location/Qualifiers					
27	1. .293962					
28	/organism="Rattus norvegicus"					
29	/mol_type="genomic DNA"					
30	/db_xref="taxon:10116"					
31	/clone="CH230-208P20"					
32	6200. .7062					
33	/note="clone_boundary"					
34	clone_end:Sp6					
35	site:					
36	end sequence:RWBKP94TVB"					
37	51702. .53500					
38	/note="wgs contig"					
39	53551. .54607					
40	/note="wgs contig"					
41	56203. .57503					
42	/note="wgs contig"					
43	174983. .176056					
44	/note="wgs contig"					
45	208600. .209401					
46	/note="clone_boundary"					



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clone_end:T7
site:
end sequence:RWBKP94TJB"
misc_feature 264125..265539
/note="wgs_end_extension
clone_end:T7"
misc_feature 267070..268441
/note="wgs_end_extension
clone_end:T7"

ORIGIN
Query Match 76.3%; Score 20.6; DB 2; Length 293962;
Best Local Similarity 85.2%; Pred. No. 19;
Matches 23; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 GTAGCCTAGTACCCCTAGTCTAGGC 27
|||||
Db 256250 GTAGCCAGCTACCTCTAGGTAGAC 256224
|||||

AC127236 188946 bp DNA linear ROD 27-NOV-2003
Mus musculus BAC clone RP24-35111 from chromosome 18, complete
sequence.
AC127236 AC127236 GI:37361086
HTG.
Mus musculus (house mouse)
ORGANISM
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE
AUTHORS Swearingen-Shahid, S.
TITLE The sequence of Mus musculus BAC clone RP24-35111
JOURNAL Unpublished (2001)
REFERENCE
AUTHORS Wilson, R.
TITLE Sequencing of Mus musculus
JOURNAL Unpublished (2001)
REFERENCE
AUTHORS McPherson, J.D. and Waterston, R.H.
TITLE Direct Submission
JOURNAL Submitted (14-JUL-2002) Genome Sequencing Center, 4444 Forest Park
Parkway, St. Louis, MO 63108, USA
REFERENCE
AUTHORS McPherson, J.D. and Waterston, R.H.
TITLE Direct Submission
JOURNAL Submitted (25-SEP-2002) Genome Sequencing Center, 4444 Forest Park
Parkway, St. Louis, MO 63108, USA
REFERENCE
AUTHORS Wilson, R.K.
TITLE Direct Submission
JOURNAL Submitted (02-OCT-2003) Genome Sequencing Center, 4444 Forest Park
Parkway, St. Louis, MO 63108, USA
REFERENCE
AUTHORS Wilson, R.
TITLE Direct Submission
JOURNAL Submitted (27-NOV-2003) Department of Genetics, Washington
University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
COMMENT On Oct 2, 2003 this sequence version replaced gi:23308121.
----- Genome Center
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu
Contact: submissions@watson.wustl.edu
----- Summary Statistics
Center project name: M_BB0351L01
-----

NOTICE: This sequence may not represent the entire insert of this
clone. It may be shorter because we only sequence overlapping
clone sections once, or longer because we provide a small overlap
```

between neighboring data submissions.

This sequence was finished as follows unless otherwise noted: all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

#### MAPPING INFORMATION:

Mapping information for this clone was provided by Dr. Wes Warren, Department of Genetics, Washington University, St. Louis MO. For additional information about the map position of this sequence, see <http://genome.wustl.edu>

#### SOURCE INFORMATION:

The RPCI-24 BAC library has been constructed by Pieter de Jong and coworkers (<http://www.chori.org>) from male C57BL/6J mouse spleen and/or brain genomic DNA. The clone and detailed information can be obtained from Pieter de Jong and coworkers at <http://www.chori.org>

#### NEIGHBORING SEQUENCE INFORMATION:

This sequence is the entire insert of the clone. This clone is overlapped by AC108434.

#### FEATURES

Location/Qualifiers	Source
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/mol_type="genomic DNA"	
/db_xref="taxon:10090"	
/chromosome="18"	
/map="18"	
/clone="RP24-35111"	
/clone_lib="RPCI-24"	
773..874	repeat_region
/rpt_family="Alu"	
920..1078	repeat_region
/rpt_family="Alu"	
1428..1643	repeat_region
/rpt_family="B2"	
1675..1845	repeat_region
/rpt_family="Alu"	
2684..2873	repeat_region
/rpt_family="B2"	
3164..3303	repeat_region
/rpt_family="Alu"	
3577..3670	repeat_region
/rpt_family="Alu"	
3688..3892	repeat_region
/rpt_family="B2"	
5876..6150	repeat_region
/rpt_family="B4"	
6252..6351	repeat_region
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6452..6680	repeat_region
/rpt_family="B4"	
6545..6681	repeat_region
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6698..6951	repeat_region
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7324..7516	repeat_region
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10777..10894	repeat_region
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11023..11190	repeat_region
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11031..11227	repeat_region
/rpt_family="B2"	
11303..11433	repeat_region
/rpt_family="Alu"	
11547..11724	repeat_region
/rpt_family="B2"	
11736..12221	repeat_region

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12388..12444
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12393..12555
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12919..13263
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13690..13828
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13846..14036
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14431..14567
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14704..14812
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16470..16651
/rpt_family="B2"
16751..16836
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16907..16994
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17163..17302
/rpt_family="B4"
17422..17450
/rpt_family="MaLR"
17451..17642
/rpt_family="B2"
complement(17566..17637)
/product="CRNA-Ser"
/Note="Likely pseudogene (HMM Sc=32.99 / Sec struct
Sc=-11.32)"
17643..17968
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18665..18753
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18685..18756
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18865..19010
/rpt_family="Alu"
19368..19574
/rpt_family="B4"
20046..20209
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20485..20641
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21092..21114
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21115..21504
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21505..21858
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21868..21995
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22114..22212
/rpt_family="Alu"
22862..23049
/rpt_family="B2"
23410..23556
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23563..23703
/rpt_family="Alu"
24073..24298
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24295..24497
/rpt_family="B2"
24498..24584
/rpt_family="L1"
24585..24732
/rpt_family="Alu"
24908..25055
/rpt_family="Alu"
25676..25782
/rpt_family="Alu"

repeat_region 25783..25991
repeat_region /rpt_family="MaLR"
26004..26094

Query Match 75.6%; Score 20.4; DB 10; Length 188946;
Best Local Similarity 95.5%; Pred. No. 24;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy 6 CTAGCTACCCCTAGGTTCTAGGC 27
||| ||||| ||||| ||||| |||||
Db 132397 CTAAGTACCCCTAGGTTCTAGGC 132418

RESULT 7
AC116089/ c 194613 bp DNA linear HTG 20-NOV-2002
LOCUS Rattus norvegicus clone CH230-344O11, WORKING DRAFT SEQUENCE, 2
DEFINITION unrounded pieces.
AC116089
VERSION AC116089.6 GI:25138296
KEYWORDS HTG; HTGS PHASE1; HTGS DRAFT; HTGS FULLTOP.
SOURCE Rattus norvegicus (Norway rat)
ORGANISM Rattus norvegicus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.
REFERENCE 1 (bases 1 to 194613)
AUTHORS Muzny, D. Marie., Metzker, M. Lee., Abramzon, S., Adams, C., Alder, J.,
Allen, C., Allen, H., Alsbrooks, S., Amin, A., Anguiano, D.,
Anyalebechi, V., Aoyagi, A., Ayodeji, M., Baca, E., Baden, H.,
Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Benahmed, F.,
Biswal, K., Blair, J., Blankenburg, K., Blyth, P., Brown, M.,
Bryant, N., Buhay, C., Burch, P., Burrell, K., Calderon, B.,
Cardenas, V., Carter, K., Cavazos, I., Cesar, H., Center, A.,
Chacko, J., Chavez, D., Chen, R., Chen, R., Chen, Y., Chen, Z., Chu, J.,
Cleveland, C., Cockrell, R., Cox, C., Coyte, M., Cree, A., D'Souza, L.,
Davila, M. L., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D.,
Delgado, O., Denson, S., Deramo, C., Ding, Y., Dinh, H., Divya, K.,
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Karpachy, S., Kelly, S., Kelly, S., Khan, Z., King, L., Kovar, C.,
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Liu, J., Liu, W., Liu, Y., London, P., Longacre, S., Lopez, J.,
Lorensu, H., Loulseg, H., Lozano, R. J., Lu, X., Ma, J.,
Maheshwari, M., Mahindartine, M., Mahmoud, M., Malloy, K., Mangum, A.,
Mangum, B., Mapua, P., Martin, K., Martin, R., Martinez, E.,
Mawhiney, S., McLeod, M. P., McNeill, T. Z., Meenen, E.,
Milosavljevic, A., Miner, G., Minja, E., Montemayor, J., Moore, S.,
Morgan, M., Morris, K., Morris, S., Munidasa, M., Murphy, M., Nair, L.,
Nankervis, C., Neal, D., Newton, N., Nguyen, N., Norris, S., Parks, K.,
Nwaokemele, O., Okwuonu, G., Olarnpunsagoon, A., Pal, S., Parks, K.,
Pasternak, S., Paul, H., Perez, A., Perez, L., Pfannkuch, C.,
Plopper, F., Poindexter, A., Popovic, D., Primus, E., Pu, L. L.,
Puazo, M., Quiroz, J., Rachlin, E., Reeves, K., Regier, M. A., Reigh, R.,
Reilly, B., Reilly, M., Ren, Y., Reuter, M., Richards, S., Riggs, F.,
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Steimle, M., Strong, R., Sutton, A., Svatek, A., Tabot, P., Taylor, C.,
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Wang, Q., Wang, S., Warren, J., Warren, R., Wei, X., White, F.,
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Wright, D., Wright, R., Wu, J., Yakub, S., Yen, J., Yoon, L., Yoon, V.,

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Yu, F., Zhang, J., Zhou, J., Zhou, X., Zhao, S., Dunn, D., von Niederhausern, A., Weiss, R., Smith, D.R., Holt, R.A., Smith, H.O., Weinstock, G. and Gibbs, R.A.  
 Direct Submission  
 Unpublished  
 2 (bases 1 to 194613)  
 Worley, K.C.  
 Submitted (24-MAR-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA  
 3 (bases 1 to 194613)  
 Direct Submission  
 Rat Genome Sequencing Consortium.  
 Submitted (20-NOV-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA  
 On Nov 20, 2002 this sequence version replaced gi:23194943.  
 The sequence in this assembly is a combination of BAC based reads and whole genome shotgun sequencing reads assembled using Atlas (<http://www.hgsc.bcm.tmc.edu/projects/rat/>). Each contig described in the feature table below represents a scaffold in the Atlas assembly (a 'contig-scaffold'). Within each contig-scaffold, individual sequence contigs are ordered and oriented, and separated by sized gaps filled with Ns to the estimated size. The sequence may extend beyond the ends of the clone and there may be sequence contigs within a contig-scaffold that consist entirely of whole genome shotgun sequence reads. Both end sequences and whole genome shotgun sequence only contigs will be indicated in the feature table.

----- Genome Center  
 Center: Baylor College of Medicine  
 Center code: BCM  
 Web site: <http://www.hgsc.bcm.tmc.edu/>  
 Contact: hgsc-help@bcm.tmc.edu  
 ----- Project Information  
 Center project name: GTCO  
 Center clone name: CH230-344011  
 ----- Summary Statistics  
 Assembly program: Phrap; version 0.990329  
 Consensus quality: 178038 bases at least Q40  
 Consensus quality: 179671 bases at least Q30  
 Consensus quality: 180674 bases at least Q20  
 Estimated insert size: 180448; sum-of-contigs estimation  
 Quality coverage: 8x in Q20 bases; sum-of-contigs estimation

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 \* NOTE: Estimated insert size may differ from sequence length  
 \* (see [http://www.hgsc.bcm.tmc.edu/docs/Genbank\\_draft\\_data.html](http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html)).  
 \* NOTE: This is a 'working draft' sequence. It currently  
 \* consists of 2 contigs. The true order of the pieces  
 \* is not known and their order in this sequence record is  
 \* arbitrary. Gaps between the contigs are represented as  
 \* runs of N, but the exact sizes of the gaps are unknown.  
 \* This record will be updated with the finished sequence  
 \* as soon as it is available and the accession number will  
 \* be preserved.

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 AC134931  
 DEFINITION  
 Oryza sativa (japonica cultivar-group) chromosome 5 clone  
 OSJNB007911, complete sequence.  
 AC134931  
 ACCESSION  
 VERSION  
 AC134931.2 GI:37360982  
 HTG.  
 KEYWORDS  
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 ORGANISM  
 Oryza sativa (japonica cultivar-group)  
 Oryza sativa (japonica cultivar-group)  
 Eukaryota; Viridiplantae; Streptophyta; Tracheophyta;  
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 Ehrhartoideae; Oryzaceae; Oryza.  
 1 (bases 1 to 124104)  
 Chow, T.-Y., Hsing, Y.-I.C., Chen, C.-S., Chen, H.-H., Liu, S.-M.,  
 Chao, Y.-T., Chang, S.-J., Chen, H.-C., Chen, S.-K., Chen, T.-R.,  
 Chen, Y.-L., Cheng, C.-H., Chung, C.-I., Han, S.-Y., Hsiao, S.-H.,  
 Hsiung, J.-N., Hsu, C.-H., Huang, J.-J., Kau, P.-I., Lee, M.-C.,  
 Leu, H.-L., Li, Y.-F., Lin, S.-J., Lin, Y.-C., Wu, S.-W., Yu, C.-Y.,  
 Yu, S.-W., Wu, H.-P. and Shaw, J.-F.  
 Oryza sativa BAC OSJNB007911 genomic sequence  
 Unpublished  
 2 (bases 1 to 124104)  
 Chow, T.-Y. and Hsing, Y.-I.C.  
 Direct Submission  
 Submitted (03-OCT-2002) Institute of Botany, Academia Sinica, 128,  
 Section 2, Academia Road, Nankang, Taipei 11529, Taiwan  
 3 (bases 1 to 124104)  
 Chow, T.-Y.  
 Direct Submission  
 Submitted (02-OCT-2003) Institute of Botany, Academia Sinica, 128,  
 Section 2, Academia Road, Nankang, Taipei 11529, Taiwan  
 4 (bases 1 to 124104)  
 Chow, T.-Y. and Hsing, Y.-I.C.  
 Direct Submission  
 Submitted (02-JUN-2004) Institute of Botany, Academia Sinica, 128,  
 Section 2, Academia Road, Nankang, Taipei 11529, Taiwan  
 On Oct 2, 2003 this sequence version replaced gi:23477780.  
 The orientation of the sequence is from Sp6 to T7 of the BAC clone.  
 Genes were predicated from the integrated results of the following:  
 BLASTN2.0, BLASTX2.0, GENSCAN (Chris Burge,  
<http://genes.mit.edu/GENSCAN.html>), Egenes  
 (<http://www.softberry.com/>), GlimmerR  
 (<http://www.tigr.org/softlab/glimmer/glimmer.html>), TWINSKAN  
 (<http://genes.cs.wustl.edu/>) and GeneSplicer  
 (<http://www.tigr.org/tdb/GeneSplicer/index.shtml>). The sequence was  
 searched against the Swiss-Prot+TrEMBL protein database, the NCBI  
 Plant EST database, the TIGR Rice Gene Index and the rice  
 full-length cDNA database (KOME,  
<http://cdna01.dna.affrc.go.jp/cDNA/>). Annotated genes are named to  
 indicate the level of evidence for their annotation. Genes with



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/translation="MSCPSCFKFKKMSKMGAREVTVVKHGASLKNSESEKLPV

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OY 3 AGCTAGTACCCCTAGGTAGGC 27
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RESULT 9
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DEFINITION Rattus norvegicus clone CH230-1016, *** SEQUENCING IN PROGRESS ***,
63 unordered pieces.
AC134515
AC134515.1 GI:23334689
VERSION    HTG; HTGS_PHASE1
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SOURCE     Rattus norvegicus
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Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.

1 (bases 1 to 119743)
Muzny, D., Marle, Metzker, M., Lee, S., Abramson, S., Adams, C., Alder, J.,
Allen, C., Allen, H., Alsbrooks, S., Amin, A., Anguiano, D.,
Anyalebechi, V., Aoyagi, A., Ayodeji, M., Baca, E., Baden, H.,
Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Benahmed, F.,
Biswal, K., Blair, J., Blankenburg, K., Blyth, P., Brown, M.,
Bryant, N., Buhay, C., Burch, P., Burrell, K., Calderon, E.,
Cardenas, V., Carter, K., Cavazos, I., Ceasar, H., Center, A.,
Chacko, J., Chavez, D., Chen, G., Chen, R., Chen, Y., Chen, Z., Chu, J.,
Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Cree, A., D'Souza, L.,
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Delgado, O., Denson, S., Deramo, C., Ding, Y., Dinh, H., Divya, K.,
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Miner, G., Minja, E., Montemayor, J., Moore, S., Morgan, M., Morris, K.,
Morris, S., Mundaesa, M., Murphy, M., Nair, L., Nankervis, C., Neal, D.,
Newton, N., Nguyen, N., Norris, S., Nwaokeme, O., Okwuonu, G.,
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Song, X.-Z., Sorrelle, R., Sosa, J., Steimle, M., Strong, R., Sutton, A.,
Svatek, A., Tabor, P., Taylor, C., Taylor, T., Thomas, N., Thomas, S.,
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TITLE  
JOURNAL  
REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
COMMENT

Wooden, H., Worley, K., Wright, D., Wright, R., Wu, J., Yakub, S.,  
Yen, J., Yoon, L., Yoon, V., Yu, F., Zhang, J., Zhou, J., Zhou, X.,  
Zhao, S., Dunn, D., von Niederhausern, A., Weiss, R., Smith, D.R.,  
Holt, R.A., Smith, H.O., Weinstein, G. and Gibbs, R.A.  
Direct Submission  
Unpublished  
2 (bases 1 to 119743)  
Rat Genome Sequencing Consortium  
Submitted (27-SEP-2002) Human Genome Sequencing Center, Department  
of Molecular and Human Genetics, Baylor College of Medicine, One  
Baylor Plaza, Houston, TX 77030, USA  
----- Genome Center  
Center: Baylor College of Medicine  
Code: BCM  
Web site: <http://www.hgsc.bcm.tmc.edu/>  
Contact: [hgsc-help@bcm.tmc.edu](mailto:hgsc-help@bcm.tmc.edu)  
----- Project Information  
Center project name: GPSS  
Center clone name: CH230-1016  
----- Summary Statistics  
Sequencing vector: Plasmid;  
Chemistry: Dye-terminator Big Dye; 100% of reads  
Assembly program: Phrap; version 0.990329  
Consensus quality: 65609 bases at least Q40  
Consensus quality: 70531 bases at least Q30  
Consensus quality: 73895 bases at least Q20

\*\*\* NOTE: Estimated insert size may differ from sequence length  
\* (see [http://www.hgsc.bcm.tmc.edu/docs/Genbank\\_draft\\_data.html](http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html)).  
\* NOTE: This is a 'working draft' sequence. It currently  
\* consists of 63 contigs. The true order of the pieces  
\* is not known and their order in this sequence record is  
\* arbitrary. Gaps between the contigs are represented as  
\* runs of N, but the exact sizes of the gaps are unknown.  
\* This record will be updated with the finished sequence  
\* as soon as it is available and the accession number will  
\* be preserved.

1 1021: contig of 1021 bp in length  
1022 1121: gap of unknown length  
1122 2410: contig of 1289 bp in length  
2411 2511: gap of unknown length  
2511 3615: contig of 1105 bp in length  
3616 3715: gap of unknown length  
3716 5208: contig of 1493 bp in length  
5209 5308: gap of unknown length  
5309 6348: contig of 1040 bp in length  
6349 6449: gap of unknown length  
6449 7557: gap of unknown length  
7557 8690: contig of 1034 bp in length  
8691 8790: gap of unknown length  
8791 9582: contig of 1192 bp in length  
9583 10082: gap of unknown length  
10083 11108: contig of 1026 bp in length  
11109 11209: gap of unknown length  
11209 12381: contig of 1173 bp in length  
12382 12481: gap of unknown length  
12482 13444: contig of 1463 bp in length  
13445 14044: gap of unknown length  
14045 15293: contig of 1249 bp in length  
15294 15393: gap of unknown length  
15394 16836: contig of 1443 bp in length  
16837 16936: gap of unknown length  
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18368 19813: contig of 1446 bp in length  
19814 19913: gap of unknown length  
19914 21224: contig of 1311 bp in length  
21225 21324: gap of unknown length  
21325 22712: contig of 1388 bp in length  
22713 22812: gap of unknown length  
22813 23813: contig of 1019 bp in length

23832 23931: gap of unknown length  
 23932 25886: contig of 1955 bp in length  
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 31980 33446: contig of 1367 bp in length  
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Query Match

73.3%; Score 19.8; DB 2; Length 119743;

Best Local Similarity 91.3%; Pred. No. 50;  
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 DEFINITION Rattus norvegicus clone CH230-17IN16, \*\*\* SEQUENCING IN PROGRESS  
 AC103070  
 ACCESSION AC103070.6 GI:30579789  
 VERSION  
 KEYWORDS HTG; HTGS\_PHASE1; HTGS\_DRAFT; HTGS\_ENRICHED.  
 SOURCE Rattus norvegicus (Norway rat)  
 ORGANISM  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;  
 Rattus.  
 1 (bases 1 to 211465)  
 Muzny, D. Marie., Metzker, M. Lee., Abramzon, S., Adams, C., Alder, J.,  
 Allen, C., Allen, H., Alsbrooks, S., Amin, A., Anguiano, D.,  
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 Hollins, B., Howells, S., Hulyk, S., Hume, J., Idlebird, D., Jackson, A.,  
 Jackson, L., Jacob, L., Jiang, H., Johnson, B., Johnson, R., Jolivet, A.,  
 Karpathy, S., Kelly, S., Kelly, S., Khan, Z., King, L., Kovar, C., Liu, J.,  
 Kowis, C., Kraft, C. L., Lebow, H., Levan, J., Lewis, L., Li, Z., Liu, J.,  
 Liu, J., Liu, W., Liu, Y., London, P., Longacre, S., Lopez, J.,  
 Lorensu, H., Loulseghe, H., Lozano, R. J., Lu, X., Ma, J.,  
 Maheshwari, M., Mahindartne, M., Mahmoud, M., Malloy, K., Mangum, A.,  
 Mangum, B., Mapua, P., Martin, K., Martin, R., Martinez, E.,  
 Mawhney, S., McLeod, M. P., McNeill, T. Z., Meenen, E.,  
 Milosavljevic, A., Miner, G., Minja, E., Montemayor, J., Moore, S.,  
 Morgan, M., Morris, K., Morris, S., Munidasa, M., Murphy, W., Nair, L.,  
 Nankervis, C., Neal, D., Newton, N., Nguyen, N., Norris, S.,  
 Nwackelme, O., Okwono, G., Olarnpunsagoon, A., Pal, S., Parks, K.,  
 Pasternak, S., Paul, H., Perez, A., Perez, L., Pfannkuch, C.,  
 Plapper, F., Poindexter, A., Popovic, D., Primus, E., Pu, L. L.,  
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 Rives, C., Rodkey, T., Rojas, A., Rose, M., Rose, R., Ruiz, S. J.,  
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 Wang, Q., Wang, S., Warren, J., Warren, R., Wei, X., White, F.,  
 Williams, G., Willson, R., Wlecyk, R., Woodson, H., Worley, K.,  
 Wright, D., Wright, R., Wu, J., Yakub, S., Yen, J., Yoon, L., Yoon, V.,  
 Yu, F., Zhang, J., Zhou, J., Zhou, X., Zhao, S., Dunn, D., von  
 Niederhausern, A., Weiss, R., Smith, D. R., Holt, R. A., Smith, H. O.,  
 Weinstock, G. and Gibbs, R. A.  
 Direct Submission  
 TITLE  
 JOURNAL  
 Unpublished

```

REFERENCE
AUTHORS
TITLE
JOURNAL

2 (bases 1 to 211465)
Worley, K.C.
Direct Submission
Submitted (24-NOV-2001) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 211465)
Rat Genome Sequencing Consortium.
Direct Submission
Submitted (13-MAY-2003) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA

COMMENT
On May 13, 2003 this sequence version replaced gi:23321751.
The sequence in this assembly is a combination of BAC based reads
and whole genome shotgun sequencing reads assembled using Atlas
(http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described
in the feature table below represents a scaffold in the Atlas
assembly (a 'contig-scaffold'). Within each contig-scaffold,
individual sequence contigs are ordered and oriented, and separated
by sized gaps filled with Ns to the estimated size. The sequence
may extend beyond the ends of the clone and there may be sequence
contigs within a contig-scaffold that consist entirely of whole
genome shotgun sequence reads. Both end sequences and whole genome
shotgun sequence only contigs will be indicated in the feature
table.

----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: GJDL
Center clone name: CH230-171N16
----- Summary Statistics
Assembly program: Atlas 3.0;
Consensus quality: 165184 bases at least Q40
Consensus quality: 169724 bases at least Q30
Consensus quality: 173497 bases at least Q20
Estimated insert size: 175743; sum-of-contigs estimation
Quality coverage: 5x in Q20 bases; sum-of-contigs estimation

* NOTE: Estimated insert size may differ from sequence length
* (see http://www.hgsc.bcm.tmc.edu/docs/genbank_draft_data.html).
* NOTE: This is a 'working draft' sequence. It currently
* consists of 2 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
* 1 15062: contig of 15062 bp in length
* 15063 15162: gap of unknown length
* 15163 211465: contig of 196303 bp in length.

FEATURES
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Location/Qualifiers
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/mol_type="genomic DNA"
/db_xref="taxon:10116"
/clone="CH230-171N16"
9750. 11143
/notes="wgs contig"
31208. 31466
/notes="wgs contig"
136133. 137193
/notes="wgs contig"
143435. 145324
/notes="wgs contig"
184822. 187065
/notes="wgs contig"
189375. 189980
/notes="clone boundary
clone_end:Sp6

misc_feature
misc_feature
misc_feature
misc_feature
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site:BCORI
end sequence: BH289014"
209353. 211465
/notes="wgs_end_extension
clone_end:Sp6"

misc_feature
Query Match 73.3%; Score 19.8; DB 2; Length 211465;
Best Local Similarity 91.3%; Pred. No. 48;
Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

ORIGIN
QY 5 CCTAGTACCCCTAGCTCTAGGC 27
|||||
Db 22022 CCTAAGTACCCCTAGGACTAGGC 22044
|||||

RESULT 11
AC095896
LOCUS
DEFINITION
AC095896 212559 bp DNA linear HTG 13-NOV-2002
Rattus norvegicus clone CH230-10N20, *** SEQUENCING IN PROGRESS
*** 10 unordered pieces.
AC095896
HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_ENRICHED.
VERSION
Rattus norvegicus (Norway rat)
KEYWORDS
SOURCE
ORGANISM
Rattus norvegicus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.
1 (bases 1 to 212559)
Muzny, D. Marie., Metzker, M. Lee., Abramson, S., Adams, C., Alder, J.,
Allen, C., Allen, H., Alsbrooks, S., Amin, A., Anguiano, D.,
Anyalebechi, V., Ayodeji, A., Ayodeji, M., Baca, E., Baden, H.,
Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Benahmed, F.,
Biswal, O., Blair, J., Blankenburg, K., Blyth, P., Brown, M.,
Bryant, N., Buhay, C., Burch, P., Burrell, K., Calderon, E.,
Cardenas, V., Carter, K., Cavazos, I., Ceasar, H., Center, A.,
Chacko, J., Chavez, D., Chen, G., Chen, Y., Chen, Y., Chen, Z., Chu, J.,
Cleaveland, C., Cockrell, R., Cox, C., Coyle, M., Cree, A., D'Souza, L.,
Davila, M. L., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D.,
Delgado, O., Denson, S., Deramo, C., Ding, Y., Dinh, H., Divya, K.,
Draper, H., Dugan-Rocha, S., Dunn, A., Durbin, K., Duval, B., Eaves, K.,
Egan, A., Escotto, M., Eugene, C., Evans, C. A., Falls, T., Fan, G.,
Fernandez, S., Finley, M., Flagg, N., Forbes, L., Foster, M., Foster, P.,
Fraser, C. M., Gabisi, A., Ganta, R., Garcia, A., Garner, T., Garza, M.,
Gebregorgis, E., Geer, K., Gill, R., Grady, M., Guerra, W., Guevara, W.,
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Puazo, M., Quiroz, J., Rachlin, E., Reeves, K., Regier, M. A., Reigh, R.,
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Rives, C., Rodkey, T., Rojas, A., Rose, M., Rose, R., Ruiz, S. J.,
Sanders, W., Savary, G., Scherer, S., Scott, G., Shatsman, S., Shen, H.,
Shetty, J., Shvartsbeyn, A., Sisson, I., Sitter, C. D., Smajda, D.,
Sneed, A., Sodergren, E., Song, X. Z., Sorelle, R., Sosa, J.,
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Valas, R., Vera, V., Villasana, D., Waldron, L., Walker, B., Wang, J.,

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Wang,Q., Wang,S., Warren,J., Warren,R., Wei,X., White,F., Williams,G., Willson,R., Wleczky,R., Wooden,H., Worley,K., Wright,D., Wright,R., Wu,J., Yakub,S., Yen,J., Yoon,L., Yoon,V., Yu,F., Zhang,J., Zhou,J., Zhou,X., Zhao,S., Dunn,D., von Niederhausern,A., Weiss,R., Smith,D.R., Holt,R.A., Smith,H.O., Weinstock,G. and Gibbs,R.A.

Direct Submission  
Unpublished  
2 (bases 1 to 212559)  
Worley,K.C.

Direct Submission  
Submitted (17-SEP-2001) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA  
3 (bases 1 to 212559)  
Rat Genome Sequencing Consortium.

Direct Submission  
Submitted (13-NOV-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA

On Nov 13, 2002 this sequence version replaced gi:23269609. The sequence in this assembly is a combination of BAC based reads and whole genome shotgun sequencing reads assembled using Atlas (<http://www.hgsc.bcm.tmc.edu/projects/rat/>). Each contig described in the feature table below represents a scaffold in the Atlas assembly (a 'contig-scaffold'). Within each contig-scaffold, individual sequence contigs are ordered and oriented, and separated by sized gaps filled with Ns to the estimated size. The sequence may extend beyond the ends of the clone and there may be sequence contigs within a contig-scaffold that consist entirely of whole genome shotgun sequence reads. Both end sequences and whole genome shotgun sequence only contigs will be indicated in the feature table.

----- Genome Center  
Center: Baylor College of Medicine  
Center code: BCM  
Web site: <http://www.hgsc.bcm.tmc.edu/>  
Contact: [hgsc-help@bcm.tmc.edu](mailto:hgsc-help@bcm.tmc.edu)

----- Project Information  
Center project name: GDUL  
Center clone name: CH230-10N20  
----- Summary Statistics  
Assembly program: Phrap; version 0.990329  
Consensus quality: 172337 bases at least Q40  
Consensus quality: 177428 bases at least Q30  
Consensus quality: 180480 bases at least Q20  
Estimated insert size: 178337; sum-of-contigs estimation  
Quality coverage: 5x in Q20 bases; sum-of-contigs estimation

-----  
\* NOTE: Estimated insert size may differ from sequence length (see [http://www.hgsc.bcm.tmc.edu/docs/Genbank\\_draft\\_data.html](http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html)).  
\* NOTE: This is a 'working draft' sequence. It currently consists of 10 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown.  
\* This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

1 45798: contig of 45798 bp in length  
\* 45799 45898: gap of unknown length  
\* 45899 146288: contig of 100370 bp in length  
\* 146289 146368: gap of unknown length  
\* 146369 150042: contig of 3674 bp in length  
\* 150043 150142: gap of unknown length  
\* 150143 162331: contig of 12189 bp in length  
\* 162332 162431: gap of unknown length  
\* 162432 199327: contig of 36896 bp in length  
\* 199328 199427: gap of unknown length  
\* 199428 200818: contig of 1391 bp in length  
\* 200819 200918: gap of unknown length  
\* 200919 202031: contig of 1113 bp in length  
\* 202032 202131: gap of unknown length

\* 202132 203608: contig of 1477 bp in length  
\* 203609 203708: gap of unknown length  
\* 203709 206315: contig of 2607 bp in length  
\* 206316 206415: gap of unknown length  
\* 206416 212559: contig of 6144 bp in length.

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misc\_feature  
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misc\_feature

ORIGIN  
Query Match 73.3%; Score 19.8; DB 2; Length 212559;  
Best Local Similarity 91.3%; Pred. No. 48;  
Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 5 CCTAGTACCCTAGGTCTAGGC 27  
Db 90486 CCTAACTATCCTAGGTCTAGGC 90508

RESULT 12  
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LOCUS  
DEFINITION  
ACCESSION  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM  
REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
MEDLINE  
PUBMED  
REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
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/dev stage="adult"  
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151..156  
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S.scrofa gene for apolipoprotein A1, exons 1-3.  
X69478.1 GI:1887  
apolipoprotein A-I; lecithin cholesterol acyltransferase cofactor;  
lipid binding.  
Sus scrofa (pig)  
Sus scrofa  
Sus scrofa  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.  
Mockel,B., Zinke,H., Flach,R., Weiss,B., Weiler-Guttler,H. and Gassen,H.G.  
Expression of apolipoprotein A-I in porcine brain endothelium in vitro  
J. Neurochem. 62 (2), 788-798 (1994)  
94125128  
8294940  
2 (bases 1 to 857)  
Moeckel,B.  
Direct Submission  
Submitted (27-NOV-1992) B. Moeckel, Inst. fuer Biochemie, Prof. Gassen, Technische Hochschule Darmstadt, Petersenstr. 22, 6100 Darmstadt, FRG  
Location/Qualifiers  
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/clone="PCR products"  
/tissue type="liver"  
/dev stage="adult"  
151..156  
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200..368  
intron



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PTMVFCDGMSIRIQIFEMSTQGLSPDLHWLGDSONVEELRTGLGLIAHGKT  
TQSFVEYVQDAMELAVARATMIQPELALLPSTMNCMDVKTLLTSQYLSFLA  
NTTFRGLSGIKVKGSTIVSSNNFPIWNLQYDPMKPMWRLGSGQGGRIVMDSGIW  
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DSLSFSLHSSNDTPIKFKKCCGYCIDLLEQLAEDMNFDFLIYVGDGKIYGAWKHG  
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WPLTMMGLGIFVALHTAIFLTLYEWKSRFGMTPKGRNKNVFSFSSALNVCYALLF  
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SUFFRNSMGPOQLMVMWNTSNLSDNQKYIFNDEGQNLGTQHQDIPLPFRRLP  
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## ORIGIN

Query Match 72.6%; Score 19.6; DB 4; Length 857;  
Best Local Similarity 84.6%; Pred. No. 90;  
Matches 22; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 TAGCCTAGTACCCCTAGGCTTAGGC 27  
Db 224 TACCCTAGTTCCTCCCGAGCTTAGGC 249

## RESULT 13

AK173314 5551 bp mRNA linear ROD 28-JUL-2004  
LOCUS Mus musculus mRNA for mKIAA1973 protein.  
DEFINITION AK173314  
ACCESSION AK173314  
VERSION AK173314.1 GI:50511212  
KEYWORDS FLI CDNA.  
SOURCE Mus musculus (house mouse)  
ORGANISM Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

## REFERENCE

1 Okazaki.N., Kikuno.R.F., Ohara.T., Inamoto.S., Koseki.H.,  
Hiraoka.S., Saga.Y., Seino.S., Nishimura.M., Kaisho.T., Hoshino.K.,  
Kitamura.H., Nagase.T., Ohara.O. and Koga.H.  
Prediction of the Coding Nucleotide Sequences of KIAA  
Gene: IV. The Complete Nucleotide Sequences of 500 Mouse  
KIAA-Homologous cDNAs Identified by Screening of Terminal Sequences  
of cDNA Clones Randomly Sampled from Size-Fractionated Libraries  
DNA Res. 11, 205-218 (2004)  
2 (bases 1 to 5551)

## JOURNAL

Okazaki.N., Kikuno.R.F., Nagase.T., Ohara.O. and Koga.H.  
Direct Submission  
Submitted (19-MAY-2004) Hisashi Koga, Kazusa DNA Research  
Institute, Laboratory for Genome Informatics; 2-6-7  
Kazusa-kamatori, Kisarazu, Chiba 292-0818, Japan  
(E-mail:mouse@kazusa.or.jp, Tel:81-438-52-3919, Fax:81-438-52-3918)  
The CREATE program supported by Japan science and technology  
corporation; cDNA full insert sequencing; Kazusa DNA Research  
Institute; cDNA library construction, clone selection and 5'- &  
3'-end one pass sequencing.

## FEATURES

Location/Qualifiers  
1..5551  
/organism="Mus musculus"  
/mol\_type="mRNA"  
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/gene="mKIAA1973"  
/note="CDS is predicted by in silico analysis. Start codon  
is not identified."  
/codon\_start=2  
/evidence="not experimental"  
/product="mKIAA1973 protein"  
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## gene

## CDS

## ORIGIN

Query Match 72.6%; Score 19.6; DB 10; Length 5551;  
Best Local Similarity 84.6%; Pred. No. 79;  
Matches 22; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 TAGCCTAGTACCCCTAGGCTTAGGC 27  
Db 4186 TAGCCTAGTTCCTCCCTTTGTTAGGC 4211

## RESULT 14

AL732521/c 207223 bp DNA linear ROD 16-JUL-2003  
LOCUS Mouse DNA sequence from clone RP23-134A17 on chromosome 4, complete  
DEFINITION AL732521  
ACCESSION AL732521  
VERSION AL732521.18 GI:32879627  
KEYWORDS HTG.  
SOURCE Mus musculus (house mouse)  
ORGANISM Mus musculus

## REFERENCE

1 (bases 1 to 207223)  
Wallis.J.  
Direct Submission  
Submitted (16-JUL-2003) Wellcome Trust Sanger Institute, Hinxton,  
Cambridgeshire, CB10 1SA, UK. E-mail enquiries:  
humquery@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk  
On Jul 16, 2003 this sequence version replaced gi:32480449.  
Sequence from the Mouse Genome Sequencing Consortium whole genome  
shotgun may have been used to confirm this sequence. Sequence data  
from the whole genome shotgun alone has only been used where it has  
a phred quality of at least 30.

## AUTHORS

Center: Wellcome Trust Sanger Institute  
Center code: SC  
Web site: http://www.sanger.ac.uk  
Contact: humquery@sanger.ac.uk

During sequence assembly data is compared from overlapping clones.  
Where differences are found these are annotated as variations  
together with a note of the overlapping clone name. Note that the  
variation annotation may not be found in the sequence submission  
corresponding to the overlapping clone, as we submit sequences with  
only a small overlap as described above.  
This sequence was finished as follows unless otherwise noted: all  
regions were either double-stranded or sequenced with an alternate  
chemistry or covered by high quality data (i.e., phred quality >=  
30); an attempt was made to resolve all sequencing problems, such  
as compressions and repeats; all regions were covered by at least  
one plasmid subclone or more than one M13 subclone; and the  
assembly was confirmed by restriction digest, except on the rare

occasion of the clone being a YAC.  
The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases:  
Em: EMBL; Sw: SWISSPROT; Tr: TREMBL; Wp: WORMPEP; Information on the WORMPEP database can be found at [http://www.sanger.ac.uk/Projects/C\\_elegans/wormpep](http://www.sanger.ac.uk/Projects/C_elegans/wormpep) RP23-134A17 is from the RPCI-23 Mouse BAC Library constructed by the group of Pieter de Jong.  
For further details see <http://www.chori.org/bacpac/home.htm>  
VECTOR: pBac3.6

## FEATURES

source Location/Qualifiers

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## ORIGIN

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Best Local Similarity 84.6%; Pred. NO. 61;  
Matches 22; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

OY 2 TAGCCTAGCTACCCCTAGGTCTAGGC 27

Db 117297 TAGCCTAGCTCCCTTTGTTAGGC 117272

## RESULT 15

AC126959

LOCUS

DEFINITION

AC126959 Rattus norvegicus 8 BAC CH230-10P12 (Children's Hospital Oakland Research Institute) complete sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Rattus norvegicus (Norway rat)

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;

Rattus

1 (bases 1 to 212481)

Muzny, D. Marie., Metzker, M. Lee., Abramson, S., Adams, C., Alder, J.,

Allen, C., Allen, H., Alsbrooks, S., Amin, A., Anguiano, D.,

Anyaletbechi, V., Aoyagi, A., Ayodeji, M., Baca, E., Baden, H.,

Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Benahmed, F.,

Biswal, K., Blair, J., Blankenburg, K., Blyth, P., Brown, M.,

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Cardenas, V., Carter, K., Cavazos, I., Ceasar, H., Center, A.,

Chacko, J., Chavez, D., Chen, G., Chen, R., Chen, Y., Chen, Z., Chu, J.,

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Davila, M., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D.,

Delgado, O., Denson, S., Deramo, C., Ding, Y., Dinh, H., Divya, K.,

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Holt, R. A., Smith, H. O., Weinstock, G. and Gibbs, R. A.

## Direct Submission

Unpublished

2 (bases 1 to 212481)

Worley, K. C.

Direct Submission

Submitted (11-JUL-2002)

Human Genome Sequencing Center, Department

of Molecular and Human Genetics, Baylor College of Medicine, One

Baylor Plaza, Houston, TX 77030, USA

3 (bases 1 to 212481)

Worley, K. C.

Direct Submission

Submitted (24-JUN-2004)

Human Genome Sequencing Center, Department

of Molecular and Human Genetics, Baylor College of Medicine, One

Baylor Plaza, Houston, TX 77030, USA

4 (bases 1 to 212481)

Worley, K. C.

Direct Submission

Submitted (24-JUN-2004)

Human Genome Sequencing Center, Department

of Molecular and Human Genetics, Baylor College of Medicine, One

Baylor Plaza, Houston, TX 77030, USA

On Jun 24, 2004 this sequence version replaced gi:30522732.

Sequencing is completed to a minimum standard of double strand

coverage with a minimum of 2 clones and 2 reads with no ambiguities

or 2 chemistries with a minimum of 2 clones and 3 reads with no

ambiguities. If the sequence quality does not meet this standard,

it will be indicated in the annotation.

Location/Qualifiers

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/db\_xref="taxon:10116"

/chromosome="8"

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/rpt\_family="MLT1b"

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3877..4035

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## COMMENT

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Best Local Similarity 84.6%; Pred. No. 61;
Matches 22; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 TAGCCTAGCTACCCCTAGCTTAGGC 27
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Search completed: October 13, 2005, 19:37:32  
Job time : 1488 secs

GenCore version 5.1.6  
Copyright (c) 1993 - 2005 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: October 13, 2005, 17:44:53 ; Search time 1122 Seconds  
(without alignments)  
142.454 Million cell updates/sec

Title: US-09-744-097A-76

Perfect score: 27

Sequence: 1 gtgcctagctaccctctagctctagc 27

Scoring table: IDENTITY\_NUC

Gapop 10.0 , Gapext 1.0

Searched: 4390206 seqs, 2959870667 residues

Total number of hits satisfying chosen parameters: 8780412

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Post-processing: Minimum Match 0%  
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Listing first 45 summaries

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- 5: Geneseqn2001bs:\*
- 6: Geneseqn2002as:\*
- 7: Geneseqn2002bs:\*
- 8: Geneseqn2003as:\*
- 9: Geneseqn2003bs:\*
- 10: Geneseqn2003cs:\*
- 11: Geneseqn2003ds:\*
- 12: Geneseqn2004as:\*
- 13: Geneseqn2004bs:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	27	100.0	27	3 AAA32245	Aaa32245 Green ter
2	27	100.0	27	3 AAA32253	Aaa32253 Distal sp
3	27	100.0	27	3 AAA32261	Aaa32261 Double di
4	27	100.0	27	3 AAA32262	Aaa32262 Double di
5	27	100.0	27	3 AAA32274	Aaa32274 One to tw
6	27	100.0	27	3 AAA32293	Aaa32293 Green ter
7	27	100.0	27	3 AAA32255	Aaa32255 Proximal
8	27	100.0	27	3 AAA32243	Aaa32243 Distal sp
9	27	100.0	27	3 AAA32264	Aaa32264 Proximal
10	27	100.0	27	3 AAA32241	Aaa32241 Proximal
11	27	100.0	27	3 AAA32275	Aaa32275 One to tw
12	27	100.0	27	4 AAS14192	Aas14192 GenetAG F
13	27	100.0	27	4 AAS14194	Aas14194 First-GRE
14	27	100.0	27	4 AAS14183	Aas14183 Fragment
15	27	100.0	27	4 AAS14198	Aas14198 First-GRE
16	27	100.0	27	4 AAS14191	Aas14191 First-GRE
17	27	100.0	27	4 AAS14181	Aas14181 Fragment
18	27	100.0	29	4 AAS14197	Aas14197 First-GRE
19	27	100.0	31	4 AAS14190	Aas14190 First-GRE
20	27	100.0	36	4 AAS14195	Aas14195 First-GRE

C 21	27	100.0	42	3 AAA32311	Aaa32311 Anti-sens
C 22	27	100.0	42	4 AAS14178	Aas14178 Modified
C 23	27	100.0	57	3 AAA32233	Aaa32233 Green rep
C 24	27	100.0	72	3 AAA32310	Aaa32310 Anti-sens
C 25	25	92.6	25	3 AAA32225	Aaa32225 Terminato
C 26	25	92.6	25	3 AAA32188	Aaa32188 Terminato
C 27	25	92.6	25	3 AAA32177	Aaa32177 Distal li
C 28	25	92.6	37	3 AAA32184	Aaa32184 Overlap l
C 29	25	92.6	37	3 AAA32174	Aaa32174 Overlap o
C 30	25	92.6	49	3 AAA32220	Aaa32220 Reporter
C 31	25	92.6	57	3 AAA32222	Aaa32222 Proximal
C 32	25	92.6	57	3 AAA32223	Aaa32223 Distal su
C 33	25	92.6	57	3 AAA32186	Aaa32186 Distal su
C 34	25	92.6	57	3 AAA32185	Aaa32185 Proximal
C 35	25	92.6	63	3 AAA32173	Aaa32173 Target sp
C 36	25	92.6	63	3 AAA32183	Aaa32183 Target ol
C 37	24	88.9	24	3 AAA32175	Aaa32175 Proximal
C 38	24	88.9	24	3 AAA32182	Aaa32182 Terminato
C 39	24	88.9	24	3 AAA32179	Aaa32179 Proximal
C 40	24	88.9	40	3 AAA32199	Aaa32199 Second re
C 41	24	88.9	40	3 AAA32191	Aaa32191 Forward r
C 42	24	88.9	40	3 AAA32193	Aaa32193 Reverse r
C 43	24	88.9	40	3 AAA32204	Aaa32204 Forward r
C 44	24	88.9	40	3 AAA32197	Aaa32197 Reporter
C 45	24	88.9	40	3 AAA32206	Aaa32206 Reverse r

## ALIGNMENTS

### RESULT 1

AAA32245

ID AAA32245 standard; DNA; 27 BP.

XX

AC AAA32245;

XX

DT 14-JUL-2000 (first entry)

XX

DE Green terminator sequence used in gene-tag reporter construction.

XX

KW Gene-tag reporter; detection; gene mapping; mutation identification;

KW cancer; mutant virus; human diagnostic; forensic; genetic analysis; ss.

XX

OS Synthetic.

XX

PN WO200004192-A1.

XX

PD 27-JAN-2000.

XX

PF 16-JUL-1999; 99WO-US016242.

XX

PR 17-JUL-1998; 98US-0093219P.

XX

PA (UYEM-) UNIV EMORY.

XX

PI Shafer DA;

XX

DR WPI; 2000-182448/16.

XX

PT New Gene-tag reporter for joining to nucleic acid probe, used, e.g. for

XX

PS mapping genes or mutational analysis.

XX

PS Example 12; Page 62; 164pp; English.

XX

CC This sequence is used in the construction of the gene-tag reporter of the invention. The invention relates to a gene-tag reporter for joining a probe, alone or with a second gene-tag reporter. The gene-tag reporter comprises a labelled double-stranded polynucleotide sequence having one or more linkers that comprise a single stranded sequence hybridisable to a complement but not to the target probe. Also included in the invention is a reporter array, comprising at least two gene-tag reporters linked end to end by hybridisable linkers. The reporters are used in various new methods for detecting and mapping genes; identifying mutations and

CC variant nucleic acids, e.g. detecting rare mutations such as those in  
 CC cancer cells or mutant viruses, but more generally in human diagnostics,  
 CC forensics, genetic analysis, analysis of environmental samples or foods.  
 CC The gene-tag reporters and associated probes, have a modular structure,  
 CC allowing simple and inexpensive probe design, and are able to generate a  
 CC distinctive signal, based on the mix and/or proportions of different  
 CC signal components. Many targets can be analysed simultaneously, using  
 CC many probes. Arrays of gene-tag reporters will provide signal  
 CC amplification  
 XX  
 SQ Sequence 27 BP; 5 A; 9 C; 7 G; 6 T; 0 U; 0 Other;

Query Match 100.0%; Score 27; DB 3; Length 27;  
 Best Local Similarity 100.0%; Pred. No. 0.0075;  
 Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTAGCCTAGTACCCCTAGGTCCTAGGC 27  
 Db 1 GTAGCCTAGTACCCCTAGGTCCTAGGC 27

RESULT 2  
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 ID AAA32253 standard; DNA; 27 BP.  
 XX  
 AC AAA32253;  
 XX  
 DT 14-JUL-2000 (first entry)  
 XX  
 DE Distal spacer oligomer A used in gene-tag reporter construction.  
 XX  
 KW Gene-tag reporter; detection; gene mapping; mutation identification;  
 KW cancer; mutant virus; human diagnostic; forensic; genetic analysis; ss.  
 XX  
 OS Synthetic.  
 XX  
 PN WO200004192-A1.  
 XX  
 PD 27-JAN-2000.  
 XX  
 PF 16-JUL-1999; 99WO-US016242.  
 XX  
 PR 17-JUL-1998; 98US-0093219P.  
 XX  
 PA (UYEM-) UNIV EMORY.  
 XX  
 PI Shafer DA;  
 XX  
 DR WPI; 2000-182448/16.  
 XX  
 PT New Gene-tag reporter for joining to nucleic acid probe, used, e.g. for  
 PT mapping genes or mutational analysis.  
 XX  
 PS Example 13; Page 67; 164pp; English.  
 XX

CC This sequence is used in the construction of the gene-tag reporter of the  
 CC invention. The invention relates to a gene-tag reporter for joining a  
 CC probe, alone or with a second gene-tag reporter. The gene-tag reporter  
 CC comprises a labelled double-stranded polynucleotide sequence having one  
 CC or more linkers that comprise a single stranded sequence hybridisable to  
 CC a complement but not to the target probe. Also included in the invention  
 CC is a reporter array, comprising at least two gene-tag reporters linked  
 CC end to end by hybridisable linkers. The reporters are used in various new  
 CC methods for detecting and mapping genes; identifying mutations and  
 CC variant nucleic acids, e.g. detecting rare mutations such as those in  
 CC cancer cells or mutant viruses, but more generally in human diagnostics,  
 CC forensics, genetic analysis, analysis of environmental samples or foods.  
 CC The gene-tag reporters and associated probes, have a modular structure,  
 CC allowing simple and inexpensive probe design, and are able to generate a  
 CC distinctive signal, based on the mix and/or proportions of different  
 CC signal components. Many targets can be analysed simultaneously, using  
 CC many probes. Arrays of gene-tag reporters will provide signal  
 CC amplification

XX Sequence 27 BP; 6 A; 7 C; 9 G; 5 T; 0 U; 0 Other;  
 SQ  
 Query Match 100.0%; Score 27; DB 3; Length 27;  
 Best Local Similarity 100.0%; Pred. No. 0.0075;  
 Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 GTAGCCTAGTACCCCTAGGTCCTAGGC 27  
 Db 27 GTAGCCTAGTACCCCTAGGTCCTAGGC 1

RESULT 3  
 AAA32261/c  
 ID AAA32261 standard; DNA; 27 BP.  
 XX  
 AC AAA32261;  
 XX  
 DT 14-JUL-2000 (first entry)  
 XX  
 DE Double distal spacer oligomer A used in gene-tag reporter construction.  
 XX  
 KW Gene-tag reporter; detection; gene mapping; mutation identification;  
 KW cancer; mutant virus; human diagnostic; forensic; genetic analysis; ss.  
 XX  
 OS Synthetic.  
 XX  
 PN WO200004192-A1.  
 XX  
 PD 27-JAN-2000.  
 XX  
 PF 16-JUL-1999; 99WO-US016242.  
 XX  
 PR 17-JUL-1998; 98US-0093219P.  
 XX  
 PA (UYEM-) UNIV EMORY.  
 XX  
 PI Shafer DA;  
 XX  
 DR WPI; 2000-182448/16.  
 XX  
 PT New Gene-tag reporter for joining to nucleic acid probe, used, e.g. for  
 PT mapping genes or mutational analysis.  
 XX  
 PS Example 14; Page 69; 164pp; English.  
 XX

CC This sequence is used in the construction of the gene-tag reporter of the  
 CC invention. The invention relates to a gene-tag reporter for joining a  
 CC probe, alone or with a second gene-tag reporter. The gene-tag reporter  
 CC comprises a labelled double-stranded polynucleotide sequence having one  
 CC or more linkers that comprise a single stranded sequence hybridisable to  
 CC a complement but not to the target probe. Also included in the invention  
 CC is a reporter array, comprising at least two gene-tag reporters linked  
 CC end to end by hybridisable linkers. The reporters are used in various new  
 CC methods for detecting and mapping genes; identifying mutations and  
 CC variant nucleic acids, e.g. detecting rare mutations such as those in  
 CC cancer cells or mutant viruses, but more generally in human diagnostics,  
 CC forensics, genetic analysis, analysis of environmental samples or foods.  
 CC The gene-tag reporters and associated probes, have a modular structure,  
 CC allowing simple and inexpensive probe design, and are able to generate a  
 CC distinctive signal, based on the mix and/or proportions of different  
 CC signal components. Many targets can be analysed simultaneously, using  
 CC many probes. Arrays of gene-tag reporters will provide signal  
 CC amplification

XX Sequence 27 BP; 6 A; 7 C; 9 G; 5 T; 0 U; 0 Other;  
 SQ  
 Query Match 100.0%; Score 27; DB 3; Length 27;  
 Best Local Similarity 100.0%; Pred. No. 0.0075;  
 Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 GTAGCCTAGTACCCCTAGGTCCTAGGC 27  
 Db 1 GTAGCCTAGTACCCCTAGGTCCTAGGC 1

```

Db      27 GTAGCCTAGCTACCCCTAGGCTAGGC 1
RESULT 4
AAA32262/c
ID      AAA32262 standard; DNA; 27 BP.
XX
AC      AAA32262;
XX
XX      14-JUL-2000 (first entry)
DT      14-JUL-2000 (first entry)
DE      One to two multilinker A used in gene-tag reporter construction.
XX
XX      Gene-tag reporter; detection; gene mapping; mutation identification;
KW      cancer; mutant virus; human diagnostic; forensic; genetic analysis; ss.
XX
OS      Synthetic.
XX
PN      WO200004192-A1.
XX
XX      27-JAN-2000.
PD
XX
XX      16-JUL-1999; 99WO-US016242.
PF
XX
XX      17-JUL-1998; 98US-0093219P.
PR
XX
XX      (UYEM-) UNIV EMORY.
PA
XX
XX      Shafer DA;
PI
XX
XX      WPI; 2000-182448/16.
DR
XX
XX      New Gene-tag reporter for joining to nucleic acid probe, used, e.g. for
PT      mapping genes or mutational analysis.
XX
XX      Example 15; Page 70; 164pp; English.
PS
XX
XX      This sequence is used in the construction of the gene-tag reporter of the
CC      invention. The invention relates to a gene-tag reporter for joining a
CC      probe, alone or with a second gene-tag reporter. The gene-tag reporter
CC      comprises a labelled double-stranded polynucleotide sequence having one
CC      or more linkers that comprise a single stranded sequence hybridisable to
CC      a complement but not to the target probe. Also included in the invention
CC      is a reporter array, comprising at least two gene-tag reporters linked
CC      end to end by hybridisable linkers. The reporters are used in various new
CC      methods for detecting and mapping genes; identifying mutations and
CC      cancer cells or mutant viruses, but more generally in human diagnostics,
CC      forensics, genetic analysis, analysis of environmental samples or foods.
CC      The gene-tag reporters and associated probes, have a modular structure,
CC      allowing simple and inexpensive probe design, and are able to generate a
CC      distinctive signal, based on the mix and/or proportions of different
CC      signal components. Many targets can be analysed simultaneously, using
CC      many probes. Arrays of gene-tag reporters will provide signal
CC      amplification
XX
XX      Sequence 27 BP; 6 A; 7 C; 9 G; 5 T; 0 U; 0 Other;
SQ      Query Match 100.0%; Score 27; DB 3; Length 27;
      Best Local Similarity 100.0%; Pred. No. 0.0075;
      Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 GTAGCCTAGCTACCCCTAGGCTAGGC 27
      |||||
Db      27 GTAGCCTAGCTACCCCTAGGCTAGGC 1

RESULT 6
AAA32293/c
ID      AAA32293 standard; DNA; 27 BP.
XX
AC      AAA32293;
XX
XX      14-JUL-2000 (first entry)
DT
XX
XX      Green set first linker A used in gene-tag reporter construction.
DE
XX
XX      Gene-tag reporter; detection; gene mapping; mutation identification;
KW      cancer; mutant virus; human diagnostic; forensic; genetic analysis; ss.
XX
XX      Synthetic.
XX

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PN WO200004192-A1.
PD 27-JAN-2000.
XX
XX 16-JUL-1999; 99WO-US016242.
XX
XX 17-JUL-1998; 98US-0093219P.
XX
XX (UYEM-) UNIV EMORY.
XX
XX Shafer DA;
XX
XX WPI; 2000-182448/16.
XX
XX New Gene-tag reporter for joining to nucleic acid probe, used, e.g. for
XX mapping genes or mutational analysis.
XX
XX Example 16; Page 72; 164pp; English.
XX
XX This sequence is used in the construction of the gene-tag reporter of the
XX invention. The invention relates to a gene-tag reporter for joining a
XX probe, alone or with a second gene-tag reporter. The gene-tag reporter
XX comprises a labelled double-stranded polynucleotide sequence having one
XX or more linkers that comprise a single stranded sequence hybridisable to
XX a complement but not to the target probe. Also included in the invention
XX is a reporter array, comprising at least two gene-tag reporters linked
XX end to end by hybridisable linkers. The reporters are used in various new
XX methods for detecting and mapping genes; identifying mutations and
XX variant nucleic acids, e.g. detecting rare mutations such as those in
XX cancer cells or mutant viruses, but more generally in human diagnostics,
XX forensics, genetic analysis, analysis of environmental samples or foods.
XX The gene-tag reporters and associated probes, have a modular structure,
XX allowing simple and inexpensive probe design, and are able to generate a
XX distinctive signal, based on the mix and/or proportions of different
XX signal components. Many targets can be analysed simultaneously, using
XX many probes. Arrays of gene-tag reporters will provide signal
XX amplification
XX
XX Sequence 27 BP; 6 A; 7 C; 9 G; 5 T; 0 U; 0 Other;
SQ
Query Match 100.0%; Score 27; DB 3; Length 27;
Best Local Similarity 100.0%; Pred. No. 0.0075;
Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTAGCCTAGCTACCCCTAGTCTAGGC 27
Db |||||
27 GTAGCCTAGCTACCCCTAGTCTAGGC 1

RESULT 7
AAA32255
ID AAA32255 standard; DNA; 27 BP.
XX
XX AAA32255;
XX
XX 14-JUL-2000 (first entry)
XX
XX Proximal spacer oligomer A used in gene-tag reporter construction.
XX
XX Gene-tag reporter; detection; gene mapping; mutation identification;
XX cancer; mutant virus; human diagnostic; forensic; genetic analysis; ss.
XX
XX Synthetic.
XX
XX WO200004192-A1.
XX
XX 27-JAN-2000.
XX
XX 16-JUL-1999; 99WO-US016242.
XX
XX 17-JUL-1998; 98US-0093219P.
XX
XX (UYEM-) UNIV EMORY.
XX
XX Shafer DA;
XX
XX WPI; 2000-182448/16.
XX
XX New Gene-tag reporter for joining to nucleic acid probe, used, e.g. for
XX mapping genes or mutational analysis.
XX
XX Example 16; Page 72; 164pp; English.
XX
XX This sequence is used in the construction of the gene-tag reporter of the
XX invention. The invention relates to a gene-tag reporter for joining a
XX probe, alone or with a second gene-tag reporter. The gene-tag reporter
XX comprises a labelled double-stranded polynucleotide sequence having one
XX or more linkers that comprise a single stranded sequence hybridisable to
XX a complement but not to the target probe. Also included in the invention
XX is a reporter array, comprising at least two gene-tag reporters linked
XX end to end by hybridisable linkers. The reporters are used in various new
XX methods for detecting and mapping genes; identifying mutations and
XX variant nucleic acids, e.g. detecting rare mutations such as those in
XX cancer cells or mutant viruses, but more generally in human diagnostics,
XX forensics, genetic analysis, analysis of environmental samples or foods.
XX The gene-tag reporters and associated probes, have a modular structure,
XX allowing simple and inexpensive probe design, and are able to generate a
XX distinctive signal, based on the mix and/or proportions of different
XX signal components. Many targets can be analysed simultaneously, using
XX many probes. Arrays of gene-tag reporters will provide signal
XX amplification
XX
XX Sequence 27 BP; 6 A; 7 C; 9 G; 5 T; 0 U; 0 Other;
SQ
Query Match 100.0%; Score 27; DB 3; Length 27;
Best Local Similarity 100.0%; Pred. No. 0.0075;
Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTAGCCTAGCTACCCCTAGTCTAGGC 27
Db |||||
27 GTAGCCTAGCTACCCCTAGTCTAGGC 1

RESULT 8
AAA32243/C
ID AAA32243 standard; DNA; 27 BP.
XX
XX AAA32243;
XX
XX 14-JUL-2000 (first entry)
XX
XX Distal spacer oligomer A used in gene-tag reporter construction.
XX
XX Gene-tag reporter; detection; gene mapping; mutation identification;
XX cancer; mutant virus; human diagnostic; forensic; genetic analysis; ss.
XX
XX Synthetic.
XX
XX WO200004192-A1.
XX
XX 27-JAN-2000.
XX
XX 16-JUL-1999; 99WO-US016242.
XX
XX 17-JUL-1998; 98US-0093219P.
XX
XX (UYEM-) UNIV EMORY.
XX
XX Shafer DA;
XX
XX WPI; 2000-182448/16.
XX
XX New Gene-tag reporter for joining to nucleic acid probe, used, e.g. for
XX mapping genes or mutational analysis.
XX
XX Example 12; Page 62; 164pp; English.
XX

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XX Shafer DA;
XX
XX WPI; 2000-182448/16.
XX
XX New Gene-tag reporter for joining to nucleic acid probe, used, e.g. for
XX mapping genes or mutational analysis.
XX
XX Example 13; Page 67; 164pp; English.
XX
XX This sequence is used in the construction of the gene-tag reporter of the
XX invention. The invention relates to a gene-tag reporter for joining a
XX probe, alone or with a second gene-tag reporter. The gene-tag reporter
XX comprises a labelled double-stranded polynucleotide sequence having one
XX or more linkers that comprise a single stranded sequence hybridisable to
XX a complement but not to the target probe. Also included in the invention
XX is a reporter array, comprising at least two gene-tag reporters linked
XX end to end by hybridisable linkers. The reporters are used in various new
XX methods for detecting and mapping genes; identifying mutations and
XX variant nucleic acids, e.g. detecting rare mutations such as those in
XX cancer cells or mutant viruses, but more generally in human diagnostics,
XX forensics, genetic analysis, analysis of environmental samples or foods.
XX The gene-tag reporters and associated probes, have a modular structure,
XX allowing simple and inexpensive probe design, and are able to generate a
XX distinctive signal, based on the mix and/or proportions of different
XX signal components. Many targets can be analysed simultaneously, using
XX many probes. Arrays of gene-tag reporters will provide signal
XX amplification
XX
XX Sequence 27 BP; 5 A; 9 C; 7 G; 6 T; 0 U; 0 Other;
SQ
Query Match 100.0%; Score 27; DB 3; Length 27;
Best Local Similarity 100.0%; Pred. No. 0.0075;
Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTAGCCTAGCTACCCCTAGTCTAGGC 27
Db |||||
1 GTAGCCTAGCTACCCCTAGTCTAGGC 27

RESULT 8
AAA32243/C
ID AAA32243 standard; DNA; 27 BP.
XX
XX AAA32243;
XX
XX 14-JUL-2000 (first entry)
XX
XX Distal spacer oligomer A used in gene-tag reporter construction.
XX
XX Gene-tag reporter; detection; gene mapping; mutation identification;
XX cancer; mutant virus; human diagnostic; forensic; genetic analysis; ss.
XX
XX Synthetic.
XX
XX WO200004192-A1.
XX
XX 27-JAN-2000.
XX
XX 16-JUL-1999; 99WO-US016242.
XX
XX 17-JUL-1998; 98US-0093219P.
XX
XX (UYEM-) UNIV EMORY.
XX
XX Shafer DA;
XX
XX WPI; 2000-182448/16.
XX
XX New Gene-tag reporter for joining to nucleic acid probe, used, e.g. for
XX mapping genes or mutational analysis.
XX
XX Example 12; Page 62; 164pp; English.
XX

```

xx This sequence is used in the construction of the gene-tag reporter of the  
 CC invention. The invention relates to a gene-tag reporter for joining a  
 CC probe, alone or with a second gene-tag reporter. The gene-tag reporter  
 CC comprises a labelled double-stranded polynucleotide sequence having one  
 CC or more linkers that comprise a single stranded sequence hybridisable to  
 CC a complement but not to the target probe. Also included in the invention  
 CC is a reporter array, comprising at least two gene-tag reporters linked  
 CC end to end by hybridisable linkers. The reporters are used in various new  
 CC methods for detecting and mapping genes; identifying mutations and  
 CC variant nucleic acids, e.g. detecting rare mutations such as those in  
 CC cancer cells or mutant viruses, but more generally in human diagnostics,  
 CC forensics, genetic analysis, analysis of environmental samples or foods.  
 CC The gene-tag reporters and associated probes, have a modular structure,  
 CC allowing simple and inexpensive probe design, and are able to generate a  
 CC distinctive signal, based on the mix and/or proportions of different  
 CC signal components. Many targets can be analysed simultaneously, using  
 CC many probes. Arrays of gene-tag reporters will provide signal  
 CC amplification  
 CC  
 SQ Sequence 27 BP; 6 A; 7 C; 9 G; 5 T; 0 U; 0 Other;

Query Match 100.0%; Score 27; DB 3; Length 27;  
 Best Local Similarity 100.0%; Pred. No. 0.0075;  
 Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTAGCCTAGCTACCCCTAGGTCTAGGC 27  
 DB 27 GTAGCCTAGCTACCCCTAGGTCTAGGC 1

RESULT 9  
 AAA32264  
 ID AAA32264 standard; DNA; 27 BP.  
 AC AAA32264;  
 XX  
 XX 14-JUL-2000 (first entry)  
 DE Proximal spacer oligomer A used in gene-tag reporter construction.  
 KW Gene-tag reporter; detection; gene mapping; mutation identification;  
 KW cancer; mutant virus; human diagnostic; forensic; genetic analysis; ss.  
 XX Synthetic.  
 OS  
 XX WO200004192-A1.  
 XX  
 XX 27-JAN-2000.  
 XX  
 PF 16-JUL-1999; 99WO-US016242.  
 XX  
 PR 17-JUL-1998; 98US-0093219P.  
 XX  
 PA (UYEM-) UNIV EMORY.  
 XX  
 PI Shafer DA;  
 XX  
 DR WPI; 2000-182448/16.  
 XX  
 PT New Gene-tag reporter for joining to nucleic acid probe, used, e.g. for  
 PT mapping genes or mutational analysis.  
 XX  
 PS Example 14; Page 69; 164pp; English.

xx This sequence is used in the construction of the gene-tag reporter of the  
 CC invention. The invention relates to a gene-tag reporter for joining a  
 CC probe, alone or with a second gene-tag reporter. The gene-tag reporter  
 CC comprises a labelled double-stranded polynucleotide sequence having one  
 CC or more linkers that comprise a single stranded sequence hybridisable to  
 CC a complement but not to the target probe. Also included in the invention  
 CC is a reporter array, comprising at least two gene-tag reporters linked  
 CC end to end by hybridisable linkers. The reporters are used in various new

CC methods for detecting and mapping genes; identifying mutations and  
 CC variant nucleic acids, e.g. detecting rare mutations such as those in  
 CC cancer cells or mutant viruses, but more generally in human diagnostics,  
 CC forensics, genetic analysis, analysis of environmental samples or foods.  
 CC The gene-tag reporters and associated probes, have a modular structure,  
 CC allowing simple and inexpensive probe design, and are able to generate a  
 CC distinctive signal, based on the mix and/or proportions of different  
 CC signal components. Many targets can be analysed simultaneously, using  
 CC many probes. Arrays of gene-tag reporters will provide signal  
 CC amplification  
 CC  
 SQ Sequence 27 BP; 5 A; 9 C; 7 G; 6 T; 0 U; 0 Other;

Query Match 100.0%; Score 27; DB 3; Length 27;  
 Best Local Similarity 100.0%; Pred. No. 0.0075;  
 Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTAGCCTAGCTACCCCTAGGTCTAGGC 27  
 DB 1 GTAGCCTAGCTACCCCTAGGTCTAGGC 27

RESULT 10  
 AAA32241  
 ID AAA32241 standard; DNA; 27 BP.  
 AC AAA32241;  
 XX  
 XX 14-JUL-2000 (first entry)  
 DE Proximal spacer oligomer A used in gene-tag reporter construction.  
 KW Gene-tag reporter; detection; gene mapping; mutation identification;  
 KW cancer; mutant virus; human diagnostic; forensic; genetic analysis; ss.  
 XX Synthetic.  
 OS  
 XX WO200004192-A1.  
 XX  
 XX 27-JAN-2000.  
 XX  
 PF 16-JUL-1999; 99WO-US016242.  
 XX  
 PR 17-JUL-1998; 98US-0093219P.  
 XX  
 PA (UYEM-) UNIV EMORY.  
 XX  
 PI Shafer DA;  
 XX  
 DR WPI; 2000-182448/16.  
 XX  
 PT New Gene-tag reporter for joining to nucleic acid probe, used, e.g. for  
 PT mapping genes or mutational analysis.  
 XX  
 PS Example 12; Page 62; 164pp; English.

xx This sequence is used in the construction of the gene-tag reporter of the  
 CC invention. The invention relates to a gene-tag reporter for joining a  
 CC probe, alone or with a second gene-tag reporter. The gene-tag reporter  
 CC comprises a labelled double-stranded polynucleotide sequence having one  
 CC or more linkers that comprise a single stranded sequence hybridisable to  
 CC a complement but not to the target probe. Also included in the invention  
 CC is a reporter array, comprising at least two gene-tag reporters linked  
 CC end to end by hybridisable linkers. The reporters are used in various new  
 CC methods for detecting and mapping genes; identifying mutations and  
 CC variant nucleic acids, e.g. detecting rare mutations such as those in  
 CC cancer cells or mutant viruses, but more generally in human diagnostics,  
 CC forensics, genetic analysis, analysis of environmental samples or foods.  
 CC The gene-tag reporters and associated probes, have a modular structure,  
 CC allowing simple and inexpensive probe design, and are able to generate a  
 CC distinctive signal, based on the mix and/or proportions of different  
 CC signal components. Many targets can be analysed simultaneously, using  
 CC many probes. Arrays of gene-tag reporters will provide signal



CC amplification  
XX Sequence 27 BP; 5 A; 9 C; 7 G; 6 T; 0 U; 0 Other;  
SQ Query Match 100.0%; Score 27; DB 3; Length 27;  
Best Local Similarity 100.0%; Pred. No. 0.0075;  
Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTAGCCTAGTACCCCTAGTCTAGGC 27  
|||  
Db 1 GTAGCCTAGTACCCCTAGTCTAGGC 27

RESULT 11  
AAA32275/c  
ID AAA32275 standard; DNA; 27 BP.  
XX  
AC AAA32275;  
XX  
DT 14-JUL-2000 (first entry)  
XX  
DE One to two multilinker B used in gene-tag reporter construction.  
XX  
KW Gene-tag reporter; detection; gene mapping; mutation identification;  
KW cancer; mutant virus; human diagnostic; forensic; genetic analysis; ss.  
XX  
OS Synthetic.  
XX WO200004192-A1.  
XX  
PD 27-JAN-2000.  
XX  
PF 16-JUL-1999; 99WO-US016242.  
XX  
PR 17-JUL-1998; 98US-0093219P.  
XX  
PA (UYEM-) UNIV EMORY.  
XX  
PI Shafer DA;  
XX  
WPI; 2000-182448/16.  
XX  
PT New Gene-tag reporter for joining to nucleic acid probe, used, e.g. for  
PT mapping genes or mutational analysis.  
XX  
PS Example 15; Page 70; 164pp; English.

This sequence is used in the construction of the gene-tag reporter of the invention. The invention relates to a gene-tag reporter for joining a probe, alone or with a second gene-tag reporter. The gene-tag reporter comprises a labelled double-stranded polynucleotide sequence having one or more linkers that comprise a single stranded sequence hybridisable to a complement but not to the target probe. Also included in the invention is a reporter array, comprising at least two gene-tag reporters linked end to end by hybridisable linkers. The reporters are used in various new methods for detecting and mapping genes; identifying mutations and variant nucleic acids, e.g. detecting rare mutations such as those in cancer cells or mutant viruses, but more generally in human diagnostics, forensics, genetic analysis, analysis of environmental samples or foods. The gene-tag reporters and associated probes, have a modular structure, allowing simple and inexpensive probe design, and are able to generate a distinctive signal, based on the mix and/or proportions of different signal components. Many targets can be analysed simultaneously, using many probes. Arrays of gene-tag reporters will provide signal amplification

Sequence 27 BP; 6 A; 7 C; 9 G; 5 T; 0 U; 0 Other;  
Query Match 100.0%; Score 27; DB 3; Length 27;  
Best Local Similarity 100.0%; Pred. No. 0.0075;  
Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTAGCCTAGTACCCCTAGTCTAGGC 27  
|||  
Db 1 GTAGCCTAGTACCCCTAGTCTAGGC 27

RESULT 12  
AAS14192/c  
ID AAS14192 standard; DNA; 27 BP.  
XX  
AC AAS14192;  
XX  
DT 18-DEC-2001 (first entry)  
XX  
DE GeneTAG First-GREEN primer used in construction of probe sets.  
XX  
KW WRAP-Probe; gene expression array; global amplification; RNA array; ss;  
KW tissue microarray; drug discovery assay; reporter binding site; forensic;  
KW diagnostic; genomic analysis; universal linker; PCR primer.  
XX  
OS Synthetic.  
XX WO200166802-A1.  
XX  
PD 13-SEP-2001.  
XX  
PF 09-MAR-2001; 2001WO-US007508.  
XX  
PR 09-MAR-2000; 2000US-0187982P.  
XX  
PA (GENE-) GENETAG TECHNOLOGY INC.  
XX  
PI Shafer DA;  
XX  
WPI; 2001-596845/67.  
XX  
PT Novel probe sets with common universal linkers at one or both ends (WRAP probes) for gene expression arrays to provide global amplification of probe set and to provide common equivalent signaling regardless of length.  
XX  
PS Example 3; Page 59; 97pp; English.

The invention relates to a probe set for gene expression arrays to provide common equivalent signalling per probe and global amplification of the set. The probe set has a pool of modified cDNA probes each probe having a central target specific segment copied from a portion of a single mRNA transcript and a universal linker (a WRAP-probe) located on one or both terminal ends. The universal linker has reporter binding sites to sites to join common reporters to the probes and primer binding sites to copy and amplify the probe. The probes and reporters are useful in diagnostic or drug discovery assays for a wide range of biomedical samples, including detection of nucleic acids and gene expression profiles in human diagnostics, forensics and genomic analysis. The methods are useful for amplifying and identifying any unknown DNA fragment and also for improving sensitivity with tissue microarrays or RNA arrays. The methods improve the quantification of gene expression and allow highly improved detection of rare transcripts or very small samples. This sequence represents a GeneTAG First-GREEN primer used in the construction of probe sets

Sequence 27 BP; 6 A; 7 C; 9 G; 5 T; 0 U; 0 Other;  
Query Match 100.0%; Score 27; DB 4; Length 27;  
Best Local Similarity 100.0%; Pred. No. 0.0075;  
Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTAGCCTAGTACCCCTAGTCTAGGC 27  
|||  
Db 27 GTAGCCTAGTACCCCTAGTCTAGGC 1

RESULT 13  
AAS14194/c  
ID AAS14194 standard; DNA; 27 BP.

xx AAS14194;  
 xx 18-DEC-2001 (first entry)  
 xx First-GREEN Chiptag primer used in construction of probe sets.  
 DE WRAP-Probe; gene expression array; global amplification; RNA array; ss;  
 xx tissue microarray; drug discovery assay; reporter binding site; forensic;  
 xx diagnostic; genomic analysis; universal linker; PCR primer.  
 xx Synthetic.  
 OS WO200166802-A1.  
 PN 13-SEP-2001.  
 PD 09-MAR-2001; 2001WO-US007508.  
 PF 09-MAR-2000; 2000US-0187982P.  
 PR (GENE-) GENETAG TECHNOLOGY INC.  
 PA Shafer DA;  
 XX WPI; 2001-596845/67.  
 XX Novel probe sets with common universal linkers at one or both ends (WRAP  
 PT probes) for gene expression arrays to provide global amplification of  
 PT probe set and to provide common equivalent signaling regardless of  
 PT length.  
 XX Example 4; Page 60; 97pp; English.  
 PS The invention relates to a probe set for gene expression arrays to  
 XX provide common equivalent signaling per probe and global amplification  
 CC of the set. The probe set has a pool of modified cDNA probes, each probe  
 CC having a central target specific segment copied from a portion of a  
 CC single mRNA transcript and a universal linker (a WRAP-Probe) located on  
 CC one or both terminal ends. The universal linker has reporter binding  
 CC sites to join common reporters to the probes and primer binding sites to  
 CC copy and amplify the probe. The probes and reporters are useful in  
 CC diagnostic or drug discovery assays for a wide range of biomedical  
 CC samples, including detection of nucleic acids and gene expression  
 CC profiles in human diagnostics, forensics and genomic analysis. The  
 CC methods are useful for amplifying and identifying any unknown DNA  
 CC fragment and also for improving sensitivity with tissue microarrays or  
 CC RNA arrays. The methods improve the quantification of gene expression and  
 CC allow highly improved detection of rare transcripts or very small  
 CC samples. This sequence represents a First-GREEN Chiptag primer used in  
 CC the construction of probe sets  
 XX Sequence 27 BP; 6 A; 7 C; 9 G; 5 T; 0 U; 0 Other;  
 SQ Query Match 100.0%; Score 27; DB 4; Length 27;  
 Best Local Similarity 100.0%; Pred. No. 0.0075;  
 Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 GTAGCCTAGCTACCCCTAGTCTAGGC 27  
 Db 27 GTAGCCTAGCTACCCCTAGTCTAGGC 1  
 RESULT 14  
 AAS14183  
 ID AAS14183 standard; DNA; 27 BP.  
 AC AAS14183;  
 XX 18-DEC-2001 (first entry)  
 XX Fragment #1 of PCR primer GR-SPC-F used in construction of probe sets.  
 DE

KW WRAP-Probe; gene expression array; global amplification; RNA array; ss;  
 KW tissue microarray; drug discovery assay; reporter binding site; forensic;  
 XX diagnostic; genomic analysis; universal linker; PCR primer.  
 OS Synthetic.  
 XX WO200166802-A1.  
 PN 13-SEP-2001.  
 PD 09-MAR-2001; 2001WO-US007508.  
 PF 09-MAR-2000; 2000US-0187982P.  
 PR (GENE-) GENETAG TECHNOLOGY INC.  
 PA Shafer DA;  
 XX WPI; 2001-596845/67.  
 XX Novel probe sets with common universal linkers at one or both ends (WRAP  
 PT probes) for gene expression arrays to provide global amplification of  
 PT probe set and to provide common equivalent signaling regardless of  
 PT length.  
 XX Disclosure; Page 90; 97pp; English.  
 PS The invention relates to a probe set for gene expression arrays to  
 CC provide common equivalent signaling per probe and global amplification  
 CC of the set. The probe set has a pool of modified cDNA probes, each probe  
 CC having a central target specific segment copied from a portion of a  
 CC single mRNA transcript and a universal linker (a WRAP-Probe) located on  
 CC one or both terminal ends. The universal linker has reporter binding  
 CC sites to join common reporters to the probes and primer binding sites to  
 CC copy and amplify the probe. The probes and reporters are useful in  
 CC diagnostic or drug discovery assays for a wide range of biomedical  
 CC samples, including detection of nucleic acids and gene expression  
 CC profiles in human diagnostics, forensics and genomic analysis. The  
 CC methods are useful for amplifying and identifying any unknown DNA  
 CC fragment and also for improving sensitivity with tissue microarrays or  
 CC RNA arrays. The methods improve the quantification of gene expression and  
 CC allow highly improved detection of rare transcripts or very small  
 CC samples. This sequence represents a fragment of a PCR primer used in the  
 CC construction of probe sets  
 XX Sequence 27 BP; 5 A; 9 C; 7 G; 6 T; 0 U; 0 Other;  
 SQ Query Match 100.0%; Score 27; DB 4; Length 27;  
 Best Local Similarity 100.0%; Pred. No. 0.0075;  
 Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 GTAGCCTAGCTACCCCTAGTCTAGGC 27  
 Db 1 GTAGCCTAGCTACCCCTAGTCTAGGC 27  
 RESULT 15  
 AAS14198  
 ID AAS14198 standard; DNA; 27 BP.  
 XX AAS14198;  
 AC 18-DEC-2001 (first entry)  
 XX First-GREEN random adapter part 2 used in construction of probe sets.  
 DE WRAP-Probe; gene expression array; global amplification; RNA array; ss;  
 KW tissue microarray; drug discovery assay; reporter binding site; forensic;  
 KW diagnostic; genomic analysis; universal linker.  
 XX Synthetic.  
 OS WO200166802-A1.

```

XX 13-SEP-2001.
XX
XX 09-MAR-2001; 2001WO-US007508.
XX
XX 09-MAR-2000; 2000US-0187982P.
XX
XX (GENE-) GENETAG TECHNOLOGY INC.
XX
XX Shafer DA;
XX
XX WPI; 2001-596845/67.
XX
XX Novel probe sets with common universal linkers at one or both ends (WRAP
XX probes) for gene expression arrays to provide global amplification of
XX probe set and to provide common equivalent signaling regardless of
XX length.
XX
XX Example 7; Page 65; 97pp; English.
XX
XX The invention relates to a probe set for gene expression arrays to
XX provide common equivalent signalling per probe and global amplification
XX of the set. The probe set has a pool of modified cDNA probes, each probe
XX having a central target specific segment copied from a portion of a
XX single mRNA transcript and a universal linker (a WRAP-Probe) located on
XX one or both terminal ends. The universal linker has reporter binding
XX sites to join common reporters to the probes and primer binding sites to
XX copy and amplify the probe. The probes and reporters are useful in
XX diagnostic or drug discovery assays for a wide range of biomedical
XX samples, including detection of nucleic acids and gene expression
XX profiles in human diagnostics, forensics and genomic analysis. The
XX methods are useful for amplifying and identifying any unknown DNA
XX fragment and also for improving sensitivity with tissue microarrays or
XX RNA arrays. The methods improve the quantification of gene expression and
XX allow highly improved detection of rare transcripts or very small
XX samples. This sequence represents a random adapter fragment used in the
XX construction of probe sets
XX
XX Sequence 27 BP; 5 A; 9 C; 7 G; 6 T; 0 U; 0 Other;
XX
XX Query Match 100.0%; Score 27; DB 4; Length 27;
XX Best Local Similarity 100.0%; Pred No. 0.0075;
XX Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX 1 GTAGCCTAGCTACCCCTAGGTTCTAGGC 27
XX |||||
XX 1 GTAGCCTAGCTACCCCTAGGTTCTAGGC 27

```

Search completed: October 13, 2005, 19:12:37  
Job time : 1123 secs

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# OM nucleic - nucleic search, using sw model

Run on: October 13, 2005, 18:25:43 ; Search time 7556 Seconds  
(without alignments)  
136.016 Million cell updates/sec

Title: US-09-744-097A-76

Perfect score: 27

Sequence: 1 gtagcctagtagtacccttaggtctagc 27

Scoring table: IDENTITY\_NUC

Gapop 10.0 , Gapext 1.0

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 68473088

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Minimum-Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database : EST.\*

1: gb\_est1.\*  
2: gb\_est2.\*  
3: gb\_est3.\*  
4: gb\_est4.\*  
5: gb\_est5.\*  
6: gb\_est6.\*  
7: gb\_est7.\*  
8: gb\_gss1.\*  
9: gb\_gss2.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
1	19.6	72.6	297	2	BB274268
2	19.6	72.6	623	1	AV325957
3	19.6	72.6	801	2	BE394439
4	19.6	72.6	896	2	BE314562
5	19.6	72.6	3378	3	AK032394
6	19.2	71.1	115	9	CG805069
7	19.2	71.1	115	9	CG805158
8	19.2	71.1	419	8	A2636851
9	19.2	71.1	453	9	CG684396
10	19.2	71.1	630	8	BZ631002
11	19.2	71.1	672	8	BZ630997
12	19.2	71.1	683	8	BZ633850
13	19.2	71.1	705	8	BH837552
14	19.2	71.1	725	8	BZ633854
15	19.2	71.1	745	9	CG351990
16	19.2	71.1	747	9	CG26350
17	19.2	71.1	828	9	CG686399
18	19.2	71.1	839	9	CG306780
19	19.2	71.1	875	9	CG227499
20	19.2	71.1	902	9	CR222205
21	19.2	71.1	951	8	CC007567
22	19	70.4	478	7	CO781043
23	19	70.4	562	8	BZ306625
24	19	70.4	774	8	CC122296

25	19	70.4	779	9	CC874542
26	19	70.4	786	9	CG063671
27	19	70.4	864	9	CG180375
28	19	70.4	865	9	CG180377
29	18.8	69.6	535	8	AZ500954
30	18.8	69.6	578	8	AZ492835
31	18.8	69.6	671	9	CR270282
32	18.8	69.6	675	9	CR273252
33	18.8	69.6	692	9	AG293534
34	18.8	69.6	711	9	AG286334
35	18.8	69.6	720	8	BZ257884
36	18.8	69.6	721	9	AG567157
37	18.6	68.9	806	7	CK316863
38	18.6	68.9	837	4	BG211849
39	18.6	68.9	869	8	BZ785030
40	18.6	68.9	880	8	BZ401569
41	18.6	68.9	954	8	CC295826
42	18.6	68.9	998	9	CG424886
43	18.6	68.9	1039	9	CC809843
44	18.2	67.4	406	8	AZ508009
45	18.2	67.4	428	6	CB097308

## ALIGNMENTS

BB274268 297 bp mRNA linear EST 07-JUL-2000  
BB274268 RIKEN full-length enriched, 10 days neonate cortex Mus  
musculus cDNA clone AB30087F07 3', mRNA sequence.

BB274268

BB274268.1 GI:8971289

EST.

Mus musculus (house mouse)

Mus musculus

Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.

1 (bases 1 to 297)

## REFERENCE

## AUTHORS

Konno, H., Aizawa, K., Akahira, S., Akiyama, J., Arakawa, T., Carninci, P., Endo, T., Fukuda, S., Fukunishi, Y., Hara, A., Hayatsu, N., Hirozane, T., Hori, F., Ishii, Y., Ishikawa, J., Ishikawa, T., Itoh, M., Izawa, M., Kadota, K., Kagawa, I., Kai, C., Kawai, J., Kikuchi, N., Kiyosawa, H., Kojima, Y., Kondo, S., Koya, S., Kurihara, C., Kusakabe, M., Matsuyama, T., Miki, R., Mizuno, Y., Nakamura, M., Oda, H., Okazaki, Y., Ono, T., Owa, C., Saito, H., Sakai, C., Sato, K., Shibata, K., Shibata, Y., Shigemoto, Y., Shinagawa, A., Shiraki, T., Sogabe, Y., Suganara, Y., Suzuki, H., Toyota, H., Tagawa, A., Takahashi, F., Tomimaga, N., Toyama, I., Tsunoda, Y., Watanabe, S., Yamamura, T., Yamana, I., Yano, R., Yasunishi, A., Yokota, T., Yoshida, K., Yoshiki, A., Yoshino, M., Muramatsu, M. and Hayashizaki, Y.

RIKEN Mouse ESTs (Konno, H., et al.)

Unpublished (2000)

Contact: Yoshihide Hayashizaki

Laboratory for Genome Exploration Research Group, RIKEN Genomic

Sciences Center (GSC), Yokohama Institute

The Institute of Physical and Chemical Research (RIKEN)

1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan

Tel: 81-45-503-9222

Fax: 81-45-503-9216

Email: genome.res@gsr.riken.jp, URL: http://genome.gsc.riken.jp/

Carninci, P., Nishiyama, Y., Westover, A., Itoh, M., Nagaoaka, S.,

Sasaki, N., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.

Thermosensitization and thermoactivation of thermolabile enzymes by

trehalose and its application for the synthesis of full length

cDNA. Proc. Natl. Acad. Sci. U.S.A. 95 (2), 520-524 (1998)

Itoh, M., Katsunari, T., Akiyama, J., Shibata, K., Izawa, M., Kawai, J.,

Tomaru, Y., Carninci, P., Shibata, Y., Ozawa, Y., Muramatsu, M.,

Okazaki, Y. and Hayashizaki, Y.

Automated filtration-based high-throughput plasmid preparation

system. Genome Res. 9 (5), 463-470 (1999)

Carninci, P. and Hayashizaki, Y.

High-efficiency full-length cDNA cloning. Methods Enzymol. 303, 19-44 (1999)  
Please visit our web site (<http://genome.rtc.riken.go.jp>) for further details.

## FEATURES

source  
1..297  
/organism="Mus musculus"  
/mol\_type="mRNA"  
/db\_xref="taxon:10090"  
/clone="A830087F07"  
/tissue\_type="cortex"  
/dev\_stage="10 days neonate"  
/lab\_host="DH10B"  
/clone\_lib="RIKEN full-length enriched, 10 days neonate cortex"  
/note="Site 1: SalI; Site 2: BamHI; cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues. 1st strand cDNA was primed with a primer [5' GAGAGAGAGATTCGAGTTAAATTAATACCCCTCCCTCCCTCC sequence [5' GAGAGAGATTCGAGTTAAATTAATACCCCTCCCTCCCTCC 3']. cDNA was cleaved with XhoI and BamHI. Vector: a modified pBluescript KS(+) after bulk excision from Lambda FLC I."

## ORIGIN

Query Match 72.6%; Score 19.6; DB 2; Length 297;  
Best Local Similarity 84.6%; Pred. No. 2.9e+02;  
Matches 22; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 TAGCCTAGCTACCCCTAGGCTAGGC 27

Db 243 TAGCCTAGCTCCCTTTGTTAGGC 268

## RESULT 2

AV325957

LOCUS AV325957 RIKEN full-length enriched, adult male medulla oblongata 623 bp mRNA linear EST 24-OCT-2001  
DEFINITION Mus musculus cDNA clone 6330407C01 3', mRNA sequence.

## ACCESSION

AV325957 GI:16395524

## VERSION

EST.

## SOURCE

Mus musculus (house mouse)

## ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

## REFERENCE

## AUTHORS

Arakawa, T., Carninci, P., Fukuda, S., Furuno, M., Hanagaki, T., Hara, A., Hiramoto, K., Hori, F., Ishii, Y., Ito, M., Kawai, J., Konno, H., Kouda, M., Koya, S., Matsuyama, T., Miyazaki, A., Nomura, K., Ohno, M., Okazaki, Y., Okido, T., Saito, R., Sakai, C., Sakai, K., Sano, H., Sasaki, D., Shibata, K., Shinagawa, A., Shiraki, T., Sogabe, Y., Suzuki, H., Tagami, M., Tagawa, A., Takahashi, F., Takeda, Y., Tanaka, T., Toyota, T., Muramatsu, M. and Hayashizaki, Y. RIKEN Mouse ESTs (Arakawa, T., et al. 2001)  
Unpublished (2001)

## TITLE

## JOURNAL

## COMMENT

On Nov 11, 1999 this sequence version replaced gi:6366009.  
Contact: Yoshihide Hayashizaki  
Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center (GSC), Yokohama Institute  
The Institute of Physical and Chemical Research (RIKEN)  
1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan  
Tel: 81-45-503-9222  
Fax: 81-45-503-9216

Email: [genome-res@gsr.riken.jp](mailto:genome-res@gsr.riken.jp), URL: <http://genome.gsc.riken.jp/>  
Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K., Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y. Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes. Genome Res. 10 (10), 1617-1630 (2000)  
wagi, K., Fujiwaki, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E., Watahiki, M., Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsura, S., Kawai, J., Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and Hayashizaki, Y.

RIKEN integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multicapillary sequencer. Genome Res. 10 (11), 1757-1771 (2000)

Konno, H., Fukunishi, Y., Shibata, K., Itoh, M., Carninci, P., Sugahara, Y. and Hayashizaki, Y.

Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)

Kondo, S., Shinagawa, A., Saito, T., Kiyosawa, H., Yamanaka, I., Aizawa, K., Fukuda, S., Hara, A., Itoh, M., Kawai, J., Shibata, K. and Hayashizaki, Y.

Computational Analysis of Full-length Mouse cDNAs Compared with Human Genome Sequences Mamm. Genome. 12, 673-677 (2001)  
Please visit our web site (<http://genome.gsc.riken.go.jp/>) for further details.

cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN.

Division of Experimental Animal Research in Riken contributed to prepare mouse tissues.

Location/Qualifiers  
1..623  
/organism="Mus musculus"  
/mol\_type="mRNA"  
/strain="C57BL/6J"  
/db\_xref="taxon:10090"  
/clone="6330407C01"  
/sex="male"  
/tissue\_type="medulla oblongata"  
/dev\_stage="adult"  
/lab\_host="DH10B"  
/clone\_lib="RIKEN full-length enriched, adult male medulla oblongata"

/note="Site 1: SalI; Site 2: BamHI; cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues. 1st strand cDNA was primed with a primer [5' GAGAGAGAGATTCGAGTTAAATTAATACCCCTCCCTCCCTCC sequence [5' GAGAGAGATTCGAGTTAAATTAATACCCCTCCCTCCCTCC 3']. cDNA was cloned into the XhoI and BamHI sites. Vector: a modified pBluescript KS(+) after bulk excision from Lambda FLC I. Cloning sites, 5' end: SalI; 3' end: BamHI"

## source

## FEATURES

Query Match 72.6%; Score 19.6; DB 1; Length 623;  
Best Local Similarity 84.6%; Pred. No. 3e+02;  
Matches 22; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

## ORIGIN

QY 2 TAGCCTAGCTACCCCTAGGCTAGGC 27

Db 569 TAGCCTAGCTCCCTTTGTTAGGC 594

## RESULT 3



MEDLINE  
PUBMED  
REFERENCE  
AUTHORS

20530913  
11076861  
4

The RIKEN Genome Exploration Research Group Phase II Team and the  
FANTOM Consortium.  
Functional annotation of a full-length mouse cDNA collection  
Nature 409, 685-690 (2001)

5  
The FANTOM Consortium and the RIKEN Genome Exploration Research  
Group Phase I & II Team.  
Analysis of the mouse transcriptome based on functional annotation  
of 60,770 full-length cDNAs  
Nature 420, 563-573 (2002)

6 (bases 1 to 3378)  
JOURNAL  
AUTHORS

Adachi, J., Aizawa, K., Akimura, T., Arakawa, T., Bono, H., Carninci, P.,  
Fukuda, S., Furuno, M., Hanagaki, T., Hara, A., Hashizume, W.,  
Hayashida, K., Hayatsu, N., Hiramoto, K., Hiraoka, T., Hirozane, T.,  
Hori, F., Imotani, K., Ishii, Y., Itoh, M., Kagawa, I., Kasukawa, T.,  
Katoh, H., Kawai, J., Kojima, Y., Kondo, S., Konno, H., Kouda, M.,  
Koya, S., Kurihara, C., Matsuyama, T., Miyazaki, A., Murata, M.,  
Nakamura, M., Nishi, K., Nomura, K., Numazaki, R., Ohno, M., Ohsato, N.,  
Okazaki, Y., Saito, R., Saitoh, H., Sakai, C., Sakai, K., Sakazume, N.,  
Sano, H., Sasaki, D., Shibata, K., Shinagawa, A., Shiraki, T.,  
Sogabe, Y., Tagami, M., Tagawa, A., Takahashi, F., Takaku-Akahira, S.,  
Takeda, Y., Tanaka, T., Tomaru, A., Toya, T., Yasunishi, A.,  
Muramatsu, M., and Hayashizaki, Y.

Direct Submission  
Submitted (16-JUL-2001) Yoshihide Hayashizaki, The Institute of  
Physical and Chemical Research (RIKEN), Laboratory for Genome  
Exploration Research Group, RIKEN Genomic Sciences Center (GSC),  
RIKEN Yokohama Institute, 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama,  
Kanagawa 230-0045, Japan (E-mail: genome-res@gsc.riken.jp,  
URL: http://genome.gsc.riken.jp/, Tel: 81-45-503-9222,  
Fax: 81-45-503-9216)

COMMENT  
cDNA library was prepared and sequenced in Mouse Genome  
Encyclopedia Project of Genome Exploration Research Group in Riken  
Genomic Sciences Center and Genome Science Laboratory in RIKEN.  
Division of Experimental Animal Research in Riken contributed to  
prepare mouse tissues.

Please visit our web site for further details.  
URL: http://genome.gsc.riken.jp/  
URL: http://fantom.gsc.riken.jp/

FEATURES  
source

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/organism="Mus musculus"  
/mol\_type="mRNA"  
/strain="C57BL/6J"  
/db\_xref="FANTOM,DB:6430537F04"  
/db\_xref="taxon:10090"  
/clone="6430537F04"  
/sex="male"  
/tissue\_type="olfactory brain"  
/clone\_lib="RIKEN full-length enriched mouse cDNA library"  
/dev\_stage="adult"

misc\_feature 1..3378  
/note="unknown EST (GB|AW494659, evidence: BLASTN, 99%,  
match=458)"

## ORIGIN

Query Match 72.6%; Score 19.6; DB 3; Length 3378;  
Best Local Similarity 84.6%; Pred. No. 3.1e+02;  
Matches 22; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Qy 2 TAGCTAGCTACCCCTAGGCTAGGC 27

Db 851 TAGCTAGCTTCCCTTTGTTAGGC 876

RESULT 6  
CG805069  
LOCUS  
DEFINITION  
1118057A09.x1 1118 - RescueMu Grid S Zea mays genomic, genomic  
survey sequence.

ACCESSION  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM

REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
COMMENT

CG805069  
CG805069.1 GI:38241110  
GSS.  
Zea mays  
Zea mays  
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD  
clade; Panicoideae; Andropogoneae; Zea.  
1 (bases 1 to 115)  
Walbot, V.  
Maize genomic sequences found using engineered RescueMu transposon  
Unpublished (2001)  
Contact: Walbot V  
Department of Biological Sciences  
Stanford University  
855 California Ave, Palo Alto, CA 94304, USA  
Tel: 650 723 2227  
Fax: 650 725 8221  
Email: walbot@stanford.edu  
Possible ligation site so sequence was trimmed. Post-ligation  
sequence submitted separately.  
Plate: 1118057 row: 10  
Class: transposon-tagged.  
Location/Qualifiers  
1..115  
/organism="Zea mays"  
/mol\_type="genomic DNA"  
/cultivar="mixed background W23/A188/B73"  
/db\_xref="taxon:4577"  
/tissue\_type="leaf"  
/dev\_stage="adult"  
/lab\_host="DH10B"  
/clone\_lib="1118 - RescueMu Grid S"  
/note="Organ: leaf; Vector: RescueMu (engineered from  
paluScript backbone); Site: 1: BamHI; Site 2: BglII;  
RescueMu is a 4.9 kb, modified maize Mu transposon  
designed to allow plasmid rescue from total genomic DNA.  
Mu elements insert preferentially into transcription  
units. For more information on RescueMu, go to the web  
site 'www.zmdb.iastate.edu' and follow the links for  
'RescueMu,' Grid S was grown at San Diego in 2002. DNA was  
extracted from leaf strips, double digested using BamHI  
and BglII, and ligated to form circular plasmids. DH10B  
cells were transformed and then screened on LB plates with  
ampicillin."

source

## ORIGIN

Query Match 71.1%; Score 19.2; DB 9; Length 115;  
Best Local Similarity 87.5%; Pred. No. 4.3e+02;  
Matches 21; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 4 GCCTAGCTACCCCTAGGCTAGGC 27

Db 58 GCCTGGCTACCCCTAGCCTAGGC 81

## RESULT 7

CG805158  
LOCUS  
DEFINITION  
1118058C04.x1 1118 - RescueMu Grid S Zea mays genomic, genomic  
survey sequence.

ACCESSION  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM

REFERENCE  
AUTHORS  
TITLE  
JOURNAL

CG805158  
CG805158.1 GI:38241278  
GSS.  
Zea mays  
Zea mays  
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD  
clade; Panicoideae; Andropogoneae; Zea.  
1 (bases 1 to 115)  
Walbot, V.  
Maize genomic sequences found using engineered RescueMu transposon  
Unpublished (2001)

## COMMENT

Contact: Walbot V  
Department of Biological Sciences  
Stanford University  
855 California Ave, Palo Alto, CA 94304, USA  
Tel: 650 723 2227  
Fax: 650 725 8221  
Email: walbot@stanford.edu

Possible ligation site so sequence was trimmed. Post-ligation  
sequence submitted separately.

Plate: 1118058 row: 10

Class: transposon-tagged

Location/Qualifiers

## FEATURES

source

1. .115

/organism="Zea mays"

/mol\_type="genomic DNA"

/cultivar="mixed background W23/A188/B73"

/db\_xref="taxon:4577"

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/dev\_stage="adult"

/lab\_host="DH10B"

/clone\_lib="1118 -- RescueMu Grid S"

/note="Organ: leaf; Vector: RescueMu (engineered from

pBlueScript backbone); Site 1: BamHI; Site 2: BglII;

RescueMu is a 4.9 kb, modified maize Mu transposon

designed to allow plasmid rescue from total genomic DNA.

Mu elements insert preferentially into transcription

units. For more information on RescueMu, go to the web

site 'www.zmdb.iastate.edu' and follow the links for

'RescueMu'. Grid S was grown at San Diego in 2002. DNA was

extracted from leaf strips, double digested using BamHI

and BglII, and ligated to form circular plasmids. DH10B

cells were transformed and then screened on LB plates with

ampicillin."

## ORIGIN

Query Match 71.1%; Score 19.2; DB 9; Length 115;

Best Local Similarity 87.5%; Pred. No. 4.3e+02;

Matches 21; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 4 GCCTAGCTACCCCTAGGCTAGGC 27

DB 58 GCCTGGCTACCCCTAGGCTAGGC 81

## RESULT 8

AZ636851

LOCUS

DEFINITION IM0495122R Mouse 10kb plasmid UUGC1M library Mus musculus genomic

clone UUGC1M0495122 R, genomic survey sequence.

ACCESSION AZ636851

VERSION AZ636851.1 GI:11759041

KEYWORDS GSS.

SOURCE Mus musculus (house mouse)

ORGANISM Mus musculus

REFERENCE 1 (bases 1 to 419)

AUTHORS Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamil, C.,

Islam, H., Longacre, S., Mahmood, M., Meenen, E., Pedersen, T.,

Kelly, M., Rose, R., Stokes, R., Tingey, A., von

Niederhauser, A. and Wright, D., Weiss, R.

Mouse whole genome scaffolding with paired end reads from 10kb

plasmid inserts

Unpublished (2000)

Contact: Robert B. Weiss

University of Utah Genome Center

Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT

84112, USA

Tel: 801 585 5606

Fax: 801 585 7177

Email: ddunn@genetics.utah.edu

Insert Length: 10000 Std Error: 0.00

## FEATURES

source

1. .419

/organism="Mus musculus"

/mol\_type="genomic DNA"

/strain="C57BL/6J"

/db\_xref="taxon:10090"

/clone="UUGC1M0495122"

/sex="Male"

/lab\_host="E. Coli strain XL10-Gold, Tl-resistant, F-"

/clone\_lib="Mouse 10kb plasmid UUGC1M library"

/note="Vector: PWD42nv; Purified genomic DNA from M.

musculus C57BL/6J (male) was obtained from the Jackson

Laboratory Mouse DNA Resource

(http://www.jax.org/resources/documents/dnares/). The DNA

was hydrodynamically sheared by repeated passage through a

0.005 inch orifice at constant velocity. The sheared DNA

was blunt end-repaired with T4 DNA polymerase and T4

polynucleotide kinase. Adaptor oligonucleotides were

ligated to the blunt ends in high molar excess. The

adapted DNA was purified and size-selected for a 9.5 to

10.5 kb range using preparative agarose gel

electrophoresis. Vector DNA was prepared from a derivative

of pMD42 (GI:4732114|gb|AF129072.1), a copy-number

inducible derivative of plasmid R1. The vector was ligated

with adaptors complementary to the insert adaptors and

purified. The sheared, adapted mouse DNA was annealed to

adapted vector DNA, and transformed into

chemically-competent E. coli XL10-Gold (Stratagene) cells

and selected for ampicillin resistance."

## ORIGIN

Query Match 71.1%; Score 19.2; DB 8; Length 419;

Best Local Similarity 87.5%; Pred. No. 4.5e+02;

Matches 21; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 3 AGCCTAGCTACCCCTAGGCTTAGG 26

DB 51 AGTCTAGCTTCCCTAGGTCACGG 74

## RESULT 9

CC684396

LOCUS

DEFINITION CC684396 453 bp DNA linear GSS 19-JUN-2003

genomic survey sequence.

ACCESSION CC684396

VERSION CC684396.1 GI:32089172

KEYWORDS GSS.

SOURCE Zea mays

ORGANISM Zea mays

REFERENCE 1 (bases 1 to 453)

AUTHORS Whitelaw, C.A., Quackenbush, J., Van Aken, S., Utterback, T.,

Resnick, A., Fraser, C.M., Budiman, M.A., Bedell, J.A., Rohlfing, T.,

Citek, R.W., Nurnberg, A., Robbins, D., and Lakey, N.

Consortium for Maize Genomics

Other GSSs: OGUKC11TH

Unpublished (2002)

Contact: Cathy Whitelaw

TIGR Medical Center Drive, Rockville, MD 20850, USA

9712 301-838-5843

Fax: 301-838-0208

Email: whitelaw@tigr.org

Seq primer: TP

Class: sheared ends.

Location/Qualifiers

## FEATURES



```

source
1. .453
/organism="Zea mays"
/mol_type="genomic DNA"
/strain="B73"
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methylation filtered genomic DNA library"

ORIGIN
Query Match 71.1%; Score 19.2; DB 9; Length 453;
Best Local Similarity 87.5%; Pred. No. 4.5e+02;
Matches 21; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 GTAGCCTAGTACCCCTAGGTCTA 24
|||||
Db 83 GTAGCCTAGTACCCCTAGCACCA 106

RESULT 10
BZ631002/c
LOCUS
DEFINITION
BZ631002 630 bp DNA linear GSS 29-JAN-2003
PAAZ280TD_ZM_0.6_1.0_KB_Zea mays genomic clone ZMMBTA007M16,
genomic survey sequence.
ACCESSION
BZ631002
VERSION
BZ631002.1 GI:28077996
KEYWORDS
GSS.
SOURCE
Zea mays
ORGANISM
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
clade; Panicoideae; Andropogoneae; Zea.
REFERENCE
1 (bases 1 to 630)
White, C.A., Quackenbush, J., Van Aken, S., Utterback, T.,
Resnick, A., Fraser, C.M., Yuan, Y., San Miguel, P., Ma, J. and
Bennetzen, J.
Maize Genomics Consortium
Unpublished (2003)
Other GSSs: PAAZ280TB
Contact: Cathy Whitelaw
TIGR
9712 Medical Center Drive, Rockville, MD 20850, USA
Tel: 301-838-5843
Fax: 301-838-0208
Email: whitelaw@tigr.org
Seq primer: TF
Class: sheared ends.
Location/Qualifiers
1. .672
/organism="Zea mays"
/mol_type="genomic DNA"
/strain="B73"
/db_xref="taxon:4577"
/clone_lib="ZMMBTA007M16"
/clone_lib="ZM_0.6_1.0_KB"
/notes="Vector: PCR4-TOPO; Site 1: EcoRI; 0.6-1.0 kb high
Cot selected genomic DNA library"

ORIGIN
Query Match 71.1%; Score 19.2; DB 8; Length 672;
Best Local Similarity 87.5%; Pred. No. 4.5e+02;
Matches 21; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 GTAGCCTAGTACCCCTAGGTCTA 24
|||||
Db 116 GTAGCCTATCCACCCCTAGGTGTA 139

RESULT 12
BZ633850
LOCUS
DEFINITION
BZ633850 683 bp DNA linear GSS 29-JAN-2003
PUAY40TB_ZM_0.6_1.0_KB_Zea mays genomic clone ZMMBTA007G07,
genomic survey sequence.
ACCESSION
BZ633850
VERSION
BZ633850.1 GI:28080844
KEYWORDS
GSS.
SOURCE
Zea mays
ORGANISM
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
clade; Panicoideae; Andropogoneae; Zea.
REFERENCE
1 (bases 1 to 683)
White, C.A., Quackenbush, J., Van Aken, S., Utterback, T.,
Resnick, A., Fraser, C.M., Yuan, Y., San Miguel, P., Ma, J. and
Bennetzen, J.
Maize Genomics Consortium
Unpublished (2003)
Other GSSs: PUAY40TD
Contact: Cathy Whitelaw
TIGR
9712 Medical Center Drive, Rockville, MD 20850, USA
Tel: 301-838-5843
Fax: 301-838-0208
Email: whitelaw@tigr.org
Seq primer: TF
Class: sheared ends.

ORIGIN
Query Match 71.1%; Score 19.2; DB 8; Length 630;
Best Local Similarity 87.5%; Pred. No. 4.5e+02;
Matches 21; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 GTAGCCTAGTACCCCTAGGTCTA 24
|||||
Db 472 GTAGCCTATCCACCCCTAGGTGTA 449

RESULT 11
BZ630997
LOCUS
DEFINITION
BZ630997 672 bp DNA linear GSS 29-JAN-2003
PAAZ280TB_ZM_0.6_1.0_KB_Zea mays genomic clone ZMMBTA007M16,
genomic survey sequence.
ACCESSION
BZ630997
VERSION
BZ630997.1 GI:28077991
KEYWORDS
GSS.
SOURCE
Zea mays
ORGANISM
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
clade; Panicoideae; Andropogoneae; Zea.
REFERENCE
1 (bases 1 to 672)
White, C.A., Quackenbush, J., Van Aken, S., Utterback, T.,
Resnick, A., Fraser, C.M., Yuan, Y., San Miguel, P., Ma, J. and
Bennetzen, J.
Maize Genomics Consortium
Unpublished (2003)
Other GSSs: PAAZ280TD
Contact: Cathy Whitelaw
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9712 Medical Center Drive, Rockville, MD 20850, USA
Tel: 301-838-5843
Fax: 301-838-0208
Email: whitelaw@tigr.org
Seq primer: TF
Class: sheared ends.

```

```

FEATURES
  source
    1..683
    /organism="Zea mays"
    /mol_type="genomic DNA"
    /strain="B73"
    /db_xref="taxon:4577"
    /clone="ZMMBTA007G07"
    /note="Vector: PCR4-TOPO; Site 1: EcoRI; 0.6-1.0 kb high
    Cot selected genomic DNA library"

ORIGIN
Query Match 71.1%; Score 19.2; DB 8; Length 683;
Best Local Similarity 87.5%; Pred. No. 4.5e+02;
Matches 21; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 GTAGCCTAGTACCCCTAGGTCTA 24
Db 116 GTAGCCTATCCACCCTAGGTGTA 139

RESULT 13
LOCUS
BH837552/c 705 bp DNA linear GSS 28-MAY-2002
DEFINITION
LMCR100001A03f Zea mays L. Zea mays genomic clone LMCRI00001A03f,
genomic survey sequence.
ACCESSION
BH837552
VERSION
BH837552.1 GI:21235430
KEYWORDS
GSS.
SOURCE
Zea mays
ORGANISM
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
clade; Panicoideae; Andropogoneae; Zea.
REFERENCE
1 (bases 1 to 705)
AUTHORS
Kim, S.W., Yu, Y., Lee, M.C., Main, D. and Wing, R.A.
TITLE
Methyl-filtration genomic sequence from maize
JOURNAL
Unpublished (2002)
COMMENT
Contact: Wing RA
Clemson University Genomics Institute
Clemson University
100 Jordan Hall, Clemson, SC 29634, USA
Tel: 864 656 7288
Fax: 864 656 4293
Email: rwing@clemson.edu
Total High Quality bases = 551
Seq primer: TAATACGACTCACTATAGGG
Class: shotgun
High quality sequence start: 12
High quality sequence stop: 690.

FEATURES
  source
    1..705
    /organism="Zea mays"
    /mol_type="genomic DNA"
    /strain="B73"
    /db_xref="taxon:4577"
    /clone="LMCR100001A03f"
    /tissue_type="Leaf"
    /lab_host="DH10B"
    /clone_lib="Zea mays L."
    /note="Vector: pGEM-T easy; Site 1: Mcr BC;
    Methyl-filtration library, Nuclei DNA was completely
    digested with Mcr BC, size fractionated and transformed
    to E.Coli.DH10B."

ORIGIN
Query Match 71.1%; Score 19.2; DB 8; Length 705;
Best Local Similarity 87.5%; Pred. No. 4.5e+02;
Matches 21; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 4 GCCTAGTACCCCTAGGTCTAGGC 27
Db 128 GCCTGGCTACCCCTAGCCCTAGGC 105

```

## RESULT 14

BZ633854/c

LOCUS

DEFINITION

ACCESSION

BZ633854

VERSION

BZ633854.1

KEYWORDS

GSS.

SOURCE

Zea mays

ORGANISM

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;

Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD

clade; Panicoideae; Andropogoneae; Zea.

REFERENCE

1 (bases 1 to 725)

AUTHORS

Whitelaw, C.A., Quackenbush, J., Van Aken, S., Utterback, T.,

Resnick, A., Fraser, C.M., Yuan, Y., San Miguel, P., Ma, J. and

Bennetzen, J.

TITLE

Maize Genomics Consortium

JOURNAL

Unpublished (2003)

COMMENT

Other\_GSSs: PUAAY40TB

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Seq primer: TF

Class: sheared ends.

Location/Qualifiers

1..725

/organism="Zea mays"

/mol\_type="genomic DNA"

/strain="B73"

/db\_xref="taxon:4577"

/clone="ZMMBTA007G07"

/clone\_lib="ZM 0.6 1.0 KB"

/note="Vector: PCR4-TOPO; Site 1: EcoRI; 0.6-1.0 kb high

Cot selected genomic DNA library"

ORIGIN

Query Match

Best Local Similarity

Matches

21; Conservative

0; Mismatches

3; Indels

0; Gaps

0;

Qy

1 GTAGCCTAGTACCCCTAGGTCTA 24

Db

567 GTAGCCTATCCACCCTAGGTGTA 544

RESULT 15

CG351990/c

LOCUS

DEFINITION

ACCESSION

CG351990

VERSION

CG351990.1

KEYWORDS

GSS.

SOURCE

Zea mays

ORGANISM

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;

Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD

clade; Panicoideae; Andropogoneae; Zea.

REFERENCE

1 (bases 1 to 745)

AUTHORS

Whitelaw, C.A., Quackenbush, J., Van Aken, S., Utterback, T.,

Resnick, A., Fraser, C.M., Budiman, M.A., Bedell, J.A., Rohlfing, T.,

Citek, R.W., Nunberg, A., Robbins, D. and Lakey, N.

TITLE

Consortium for Maize Genomics

JOURNAL

Unpublished (2002)

COMMENT

Other\_GSSs: OGVC84TV

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Email: whitelaw@tigr.org

Seq primer: TF

Class: sheared ends.

Location/Qualifiers

1..745

/organism="Zea mays"

/mol\_type="genomic DNA"

/strain="B73"

/db\_xref="taxon:4577"

/clone="ZMMBTA007G07"

/clone\_lib="ZM 0.6 1.0 KB"

/note="Vector: PCR4-TOPO; Site 1: EcoRI; 0.6-1.0 kb high

Cot selected genomic DNA library"

ORIGIN

Query Match

Best Local Similarity

Matches

21; Conservative

0; Mismatches

3; Indels

0; Gaps

0;

Qy

4 GCCTAGTACCCCTAGGTCTAGGC 27

Db

128 GCCTGGCTACCCCTAGCCCTAGGC 105

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 Fax: 301-838-0208  
 Email: whitelaw@tigr.org  
 Seq primer: TR  
 Class: Sheared ends.

FEATURES

source

Location/Qualifiers

1. .745  
 /organism="Zea mays"  
 /mol\_type="genomic DNA"  
 /strain="B73"  
 /db\_xref="taxon:4577"  
 /clone\_lib="ZM\_0.7-1.5\_KB"  
 /note="Vector: pBCSK-; Site\_1: HincII; 0.7-1.5 kb  
 methylation filtered genomic DNA library"

ORIGIN

Query Match 71.1%; Score 19.2; DB 9; Length 745;  
 Best Local Similarity 87.5%; Pred. No. 4.5e+02;  
 Matches 21; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 4 GCCTAGCTACCCCTAGCTAGGC 27  
 |||||  
 Db 389 GCCTGGCTACCCCTAGCCCTAGGC 366

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